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Date of Application:	December 25, 2002
Application Number:	2002-373531
[ST.10/C]:	JP2002-373531
Applicant(s):	JSR Corporation

December 11, 2003

Commissioner, Patent Office
Yasuo IMAI

Certificate No. 2003-3102673

[Document Name]

APPLICATION FOR PATENT

[Preference Number]

SRR10185

[Filing Date]

December 25, 2002

[Submitted to]

Commissioner, Patent Office

[International Patent Classification]

G03F 7/004

H01L 21/027

H01L 21/30

C07C317/12

C07C317/14

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[Designation of Fees]

[Advanced Payment Registration Number] 044428

[Amount Paid] 21,000 Yen

[List of Appended Documents]

[Document Name] Specification 1

[Document Name] Drawings 1

[Document Name] Abstract 1

[General Power of Attorney No.] 9116687

[Proof Requirement] Requested

HISTORICAL INFORMATION ON APPLICANT

Identification No.

[000004178]

1. Date of Alteration

December 10, 1997

[Reason for Alteration]

Alteration of Name

Address:

2-11-24, Tsukiji, Chuo-ku, Tokyo

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2. Date of Alteration

May 6, 2003

[Reason for Alteration]

Alteration of Address

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3. Date of Alteration

September 1, 2003

[Reason for Alteration]

Alteration of Name

Address:

5-6-10, Tsukiji, Chuo-ku, Tokyo

Name:

JSR Corporation

[Document Name] Specification

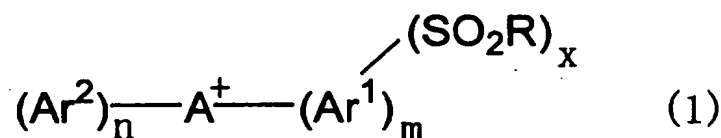
[Title of the Invention]

PHOTOACID GENERATOR, ONIUM SALT COMPOUND AND POSITIVE-TONE
RADIATION-SENSITIVE RESIN COMPOSITION

5 [Claims]

[Claim 1] A photoacid generator for photoresist comprising at least one onium
salt compound having a cation moiety of the following formula (1),

[Chemical Formula 1]



10

wherein A represents an iodine atom or a sulfur atom, when A is an iodine atom, m is 1
or 2 and n is 0 or 1, provided that (m+n) = 2, and x is an integer of 1-10, and when A is
a sulfur atom, m is 1-3 and n is 0-2, provided that (m+n) = 3, and x is an integer of 1-15;
Ar¹ and Ar² individually represent a substituted or unsubstituted monovalent aromatic

15 hydrocarbon group having 6-20 carbon atoms or a substituted or unsubstituted
monovalent heterocyclic group having 3-20 atoms, or Ar¹ and Ar² mutually bond
together with A⁺ in the formula to form a group possessing a cyclic structure with 3-8
atoms; and the x-number of -SO₂R groups bond together with one or more of the
m-number of Ar¹ groups, wherein R represents a substituted or unsubstituted alkyl

20 group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic
hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon
atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having
6-20 carbon atoms, a substituted or unsubstituted monovalent heterocyclic group having
3-20 atoms, or a -N(R')₂ group (wherein R' individually represent a hydrogen atom, a

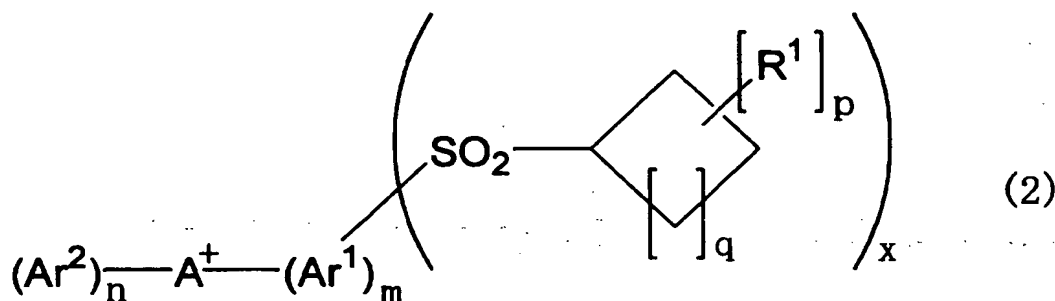
25 substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or

unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, or a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or two R' groups form, in combination and together with the nitrogen atom in the formula, a group having a cyclic structure with 3-8 atoms.)

[Claim 2] The photoacid generator for photoresist according to claim 1, wherein A in formula (1) is a sulfur atom.

[Claim 3] An onium salt compound having a cation moiety of the following formula (2),

[Chemical Formula 2]



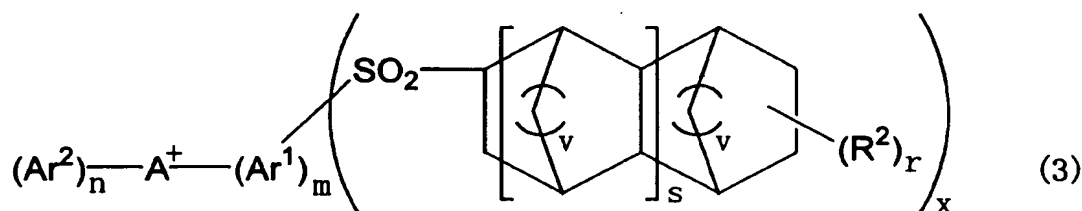
wherein A, Ar¹, m, Ar², n, and x are respectively the same as A, Ar¹, m, Ar², n, and x in the formula (1), R¹ represents a substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or a -N(R'')₂ group (wherein R'' individually represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group

having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, or a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or two R'' groups form, in combination and together with the nitrogen atom in the formula, a group having a cyclic structure with 3-8 atoms); p is an integer of 0-16, and q is an integer of 0-8.)

[Claim 4] The onium salt compound according to claim 3, wherein A in formula (2) is a sulfur atom.

[Claim 5] An onium salt compound having a cation moiety of the following formula (3),

[Chemical Formula 3]



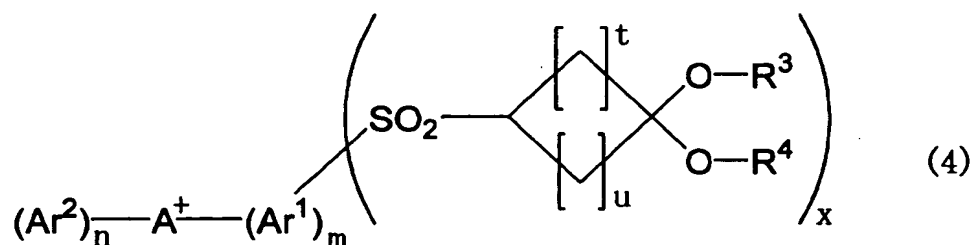
wherein A, Ar¹, m, Ar², n, and x are respectively the same as A, Ar¹, m, Ar², n, and x in the formula (1), R² represents a substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or a -N(R''')₂ group (wherein R''' individually represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, or a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or two R''' groups form, in

combination and together with the nitrogen atom in the formula, a group having a cyclic structure with 3-8 atoms); r is an integer of 0-6, s is an integer of 0-5, and v is an integer of 1-3.

[Claim 6] The onium salt compound according to claim 5, wherein A in formula (3) is a sulfur.

[Claim 7] An onium salt compound having a cation moiety of the following formula (4),

[Chemical Formula 4]



wherein A, Ar¹, m, Ar², n, and x are respectively the same as A, Ar¹, m, Ar², n, and x in the formula (1), R³ and R⁴ individually represent a substituted or unsubstituted alkyl group having 1-20 carbon atoms or a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, or R³ and R⁴ may form, in combination and together with one carbon atom and two oxygen atoms in the formula, a group having a cyclic structure with 4-10 atoms; and t and u are integers of 0-5, provided that (t+u) ≥ 1.

[Claim 8] The onium salt compound according to claim 7, wherein A in formula (4) is a sulfur.

[Claim 9] A photoacid generator obtained from any one of the onium salt compounds of claims 3-8.

[Claim 10] A positive-tone radiation-sensitive resin composition comprising: (A) a photoacid generator comprising at least one compound selected from the group

consisting of the photoacid generator for photoresist of claim 1 or claim 2 and the photoacid generator of claim 9, and (B) a resin comprising an acid-dissociable group and being insoluble or scarcely soluble in alkali, but becoming alkali soluble when the acid-dissociable group dissociates.

5 [Claim 11] The positive-tone radiation-sensitive resin composition according to claim 10, further comprising (C) an acid diffusion controller.

[Detailed Description of the Invention]

[0001]

[Technical field of the Invention]

10 The present invention relates to a photoacid generator (for photoresist) comprising an onium salt compound having a sulfone structure bonded to an aromatic hydrocarbon group or a heterocyclic group, wherein the photoacid generator is responsive to active radiations such as deep ultraviolet rays represented by a KrF excimer laser, ArF excimer laser, F₂ excimer laser, and EUV, as well as to electron
15 beams, a novel onium salt compound useful in the photoacid generator, and a positive-tone radiation-sensitive resin composition comprising the photoacid generator useful for microfabrication represented by the manufacture of integrated circuit devices.

[0002]

[Description of Background Art]

20 In the field of microfabrication represented by the manufacture of integrated circuit devices, lithographic technology enabling microfabrication with a line width of 0.30 μm or less has been demanded in order to increase the degree of integration in recent years. In a conventional lithographic process, near ultraviolet rays such as i-line radiation have been generally used. However, it is difficult to perform
25 microfabrication with a line width of sub-quarter micron using near ultraviolet rays. Therefore, in order to enable microfabrication with a line width of 0.30 μm or less, utilization of radiation with a shorter wavelength has been studied.

As radiation with a shorter wavelength, deep ultraviolet rays represented by a line spectrum of a mercury lamp and an excimer laser, X-rays, electron beams, and the like can be given. Of these, deep ultraviolet rays such as a KrF excimer laser (wavelength: 248 nm), an ArF excimer laser (wavelength: 193 nm), an F₂ excimer laser (wavelength: 157 nm), or EUV (wavelength 13 nm), and electron beams have attracted attention.

[0003]

As a radiation-sensitive resin composition applicable to short wavelength radiations, a number of compositions utilizing a chemical amplification effect between a component having an acid functional group protected with an acid-dissociable group and a photoacid generator which generates an acid upon irradiation (hereinafter referred to as "exposure") has been proposed. Such a composition is hereinafter referred to as a chemically-amplified radiation-sensitive composition. For example, Patent Document 1 discloses a chemically-amplified radiation-sensitive composition containing a polymer of which the carboxyl group or phenolic hydroxyl group is protected with a t-butoxycarbonyl group and a photoacid generator. This composition utilizes the effect of the polymer to release the t-butoxycarbonyl group by the action of an acid generated by exposure to form an acidic functional group such as a carboxyl group or a phenolic hydroxyl group, which renders an exposed area on a resist film readily soluble in an alkaline developer.

However, most of the photoacid generators currently used for conventional chemically-amplified radiation-sensitive compositions have only insufficient sensitivity. A sulfonium salt compound, which has been regarded as having the highest sensitivity, exhibits only very low base resistance, posing a serious problem of storage stability of the composition. The type of acid diffusion inhibitor, which is a basic additive for improving process stability as a resist, is also limited.

[0004]

[Patent Document 1]

Japanese Patent Application Laid-open No. 59-45439

[0005]

Therefore, development of a highly sensitive, more excellent photoacid generator exhibiting superior base resistance and excellent storage stability is strongly desired.

5 [0006]

Patent Document 2 and Non-patent Document 1 disclose that a sulfonium salt compound having a sulfonyl structure bonded with an aromatic ring is useful as a photoinitiator due to a cation mechanism. However, no sulfonium salt compound having a sulfonyl structure bonded with an alicyclic ring has been synthesized heretofore.

[0007]

[Patent Document 2]

Japanese Patent Application Laid-open No. 63-36332

[Non-patent document 1]

15 J. of Polymer Sci., Polymer Chemistry Edition, Vol. 18, pp. 2697-2714 (1980)

[0008]

[Problems to be Solved by the Invention]

The present invention has been completed in view of the above situation and has an object of providing a novel photoacid generator (for photoresist) responsive to active radiations such as deep ultraviolet rays represented by a KrF excimer laser, ArF excimer laser, F₂ excimer laser, and EUV, as well as to electron beams, having superior storage stability, and used for a radiation-sensitive resin composition as a high sensitivity, high resolution chemically-amplified photoresist. Another object of the present invention is to provide an onium salt compound suitable for a photoacid generator used as a high sensitivity and high resolution chemically-amplified photoresist, which is responsive to the above active radiations and has a superior storage stability, and a positive-tone radiation-sensitive resin composition comprising this photoacid generator (for

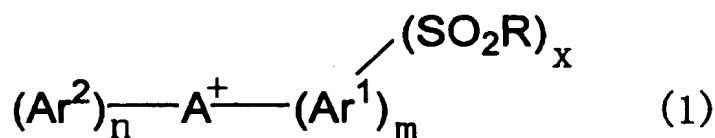
photoresist).

[0009]

First, the above object can be accomplished in the present invention by a photoacid generator for photoresist (hereinafter referred to as “photoacid generator (A1)”) comprising at least one onium salt compound having a cation moiety of the following formula (1) (hereinafter referred to as “onium salt compound (1)”),

[0010]

[Chemical Formula 5]



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[0011]

wherein A represents an iodine atom or a sulfur atom, when A is an iodine atom, m is 1 or 2 and n is 0 or 1, provided that (m+n) = 2, and x is an integer of 1-10, and when A is a sulfur atom, m is an integer of 1-3 and n is an integer of 0-2, provided that (m+n) = 3, and x is an integer of 1-15; Ar¹ and Ar² individually represent a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms or a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or Ar¹ and Ar² mutually bond together with A⁺ in the formula to form a group possessing a cyclic structure with 3-8 atoms; and the x-number of -SO₂R groups bond together with one or more of the m-number of Ar¹ groups, wherein R represents a substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or a -N(R')₂ group (wherein R'

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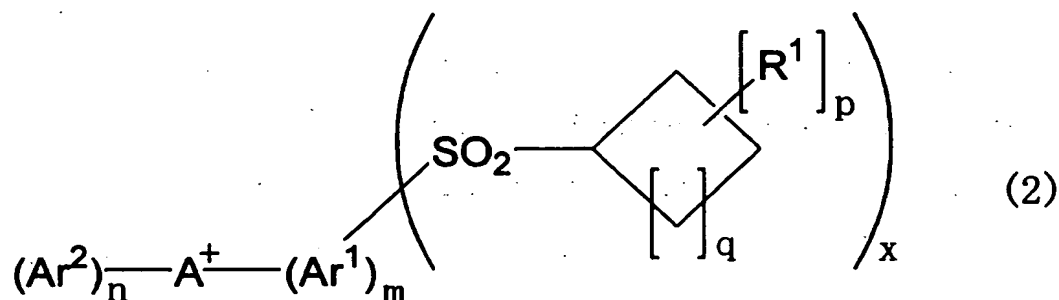
individually represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, or a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or two R' groups form, in combination and together with the nitrogen atom in the formula, a group having a cyclic structure with 3-8 atoms.)

[0012]

Second, the above object can be achieved in the present invention by an onium salt compound having a cation moiety represented by the following formula (2) (hereinafter referred to as "onium salt compound (2)"),

[0013]

[Chemical Formula 6]



[0014]

wherein A, Ar¹, m, Ar², n, and x are respectively the same as A, Ar¹, m, Ar², n, and x in the formula (1), R¹ represents a substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or a

-N(R'')₂ group (wherein R'' individually represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, or a substituted or unsubstituted monovalent heterocyclic group having 3-20 atoms, or two R'' groups form, in combination and together with the nitrogen atom in the formula, a group having a cyclic structure with 3-8 atoms); p is an integer of 0-16, and q is an integer of 0-8.

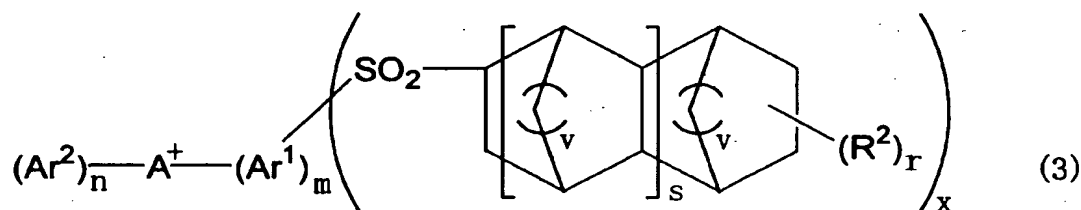
Hereinafter, the onium salt compound (2) used as the photoacid generator will be referred to from time to time as "acid generator (A2)".

[0015]

Third, the above object can be achieved in the present invention by an onium salt compound having a cation moiety represented by the following formula (3) (hereinafter referred to as "onium salt compound (3)"),

[0016]

[Chemical Formula 7]



[0017]

wherein A, Ar¹, m, Ar², n, and x are respectively the same as A, Ar¹, m, Ar², n, and x in the formula (1), R² represents a substituted or unsubstituted alkyl group having 1-20 carbon atoms, a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, an alkenyl group having 2-20 carbon atoms, a substituted or unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, a

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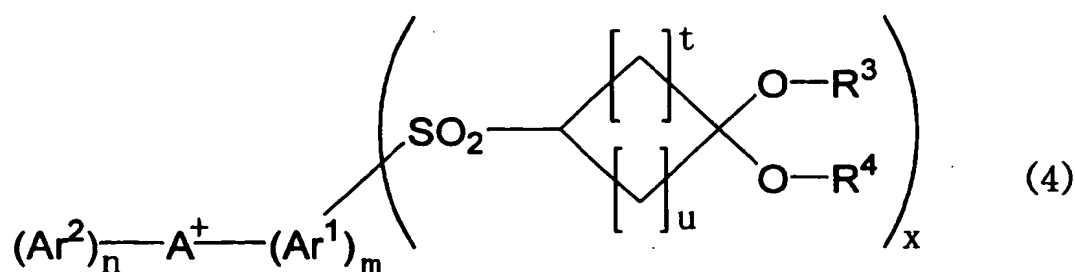
referred to from time to time as “acid generator (A3)”.

[0018]

15

[0019]

[Chemical Formula 8]



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[0020]

wherein A, Ar¹, m, Ar², n, and x are respectively the same as A, Ar¹, m, Ar², n, and x in the formula (1), R³ and R⁴ individually represent a substituted or unsubstituted alkyl

group having 1-20 carbon atoms or a substituted or unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, or R^3 and R^4 may form, in combination and together with one carbon atom and two oxygen atoms in the formula, a group having a cyclic structure with 4-10 atoms; and t and u are integers of 0-5, provided that
5 (t+u) \geq 1.

Hereinafter, the onium salt compound (4) used as the photoacid generator will be referred to from time to time as "acid generator (A4)".

[0021]

Fifth, the above objective can be achieved in the present invention by a photoacid
10 generator obtained from the onium salt compound (2), onium salt compound (3), or the onium salt compound (4).

[0022]

Sixth, the above object can be achieved in the present invention by a positive-tone radiation-sensitive resin composition comprising (A) a photoacid
15 generator containing at least one acid generator selected from the group consisting of (A1) - (A4) as an essential component and (B) an acid-dissociable group-containing resin which is insoluble or scarcely soluble in alkali, but becomes alkali soluble when the acid-dissociable group dissociates.

[0023]

20 [Preferred Embodiment of the Invention]

The present invention will be described in more detail below.

Acid generators (A1) - (A4)

The common points of the acid generator (A1) and the onium salt compounds (2) - (4) will now be explained.

25 In the formulas (1) - (4), A is preferably a sulfur atom and x is preferably 1-3.

[0024]

In the formulas (1) - (4), the number of carbon atoms in the unsubstituted

monovalent aromatic hydrocarbon group having 6-20 carbon atoms represented by Ar¹ and Ar² is preferably 6-12, more preferably 6-10, and particularly preferably 6-8.

As specific examples of the monovalent aromatic hydrocarbon group, a phenyl group, o-tolyl group, m-tolyl group, p-tolyl group, benzyl group, o-methylbenzyl group, m-methylbenzyl group, p-methylbenzyl group, 2,3-xylyl group, 2,4-xylyl group, 2,5-xylyl group, 2,6-xylyl group, 3,4-xylyl group, 5,5-xylyl group, mesityl group, 1-naphthyl group, 2-naphthyl group, 1-anthryl group, and 9-anthryl group can be given.

[0025]

The unsubstituted monovalent heterocyclic group having 3-20 atoms represented by Ar¹ and Ar² contains one or more hetero atoms such as a nitrogen atom, oxygen atom, and sulfur atom. The number of the atoms in the monovalent heterocyclic group is preferably 4-12, more preferably 5-10, and particularly preferably 5-7.

As specific examples of the monovalent heterocyclic group, 2-pyridyl group, 3-pyridyl group, 4-pyridyl group, 2-imidazolyl group, 4-imidazolyl group, 5-imidazolyl group, 2-morpholinyl group, 3-morpholinyl group, 2-piperidinyl group, 3-piperidinyl group, 4-piperidinyl group, 2-pyrrolidinyl group, and 3-pyrrolidinyl group can be given.

[0026]

As examples of the substituents for the substituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms and the substituted monovalent heterocyclic group having 3-20 atoms represented by Ar¹ and Ar², a hydroxyl group, a carboxyl group, an oxo group (=O), an alkyl group having 1-4 carbon atoms (such as a methyl group, ethyl group, n-propyl group, i-propyl group, n-butyl group, and t-butyl group), a hydroxyalkyl group having 1-4 carbon atoms (such as a hydroxymethyl group, 1-hydroxyethyl group, 2-hydroxyethyl group, 1-hydroxypropyl group, 2-hydroxypropyl group, 3-hydroxypropyl group, 1-hydroxybutyl group, 2-hydroxybutyl group, 3-hydroxybutyl group, and 4-hydroxybutyl group), an alkoxyl group having 1-4 carbon atoms (such as a methoxy group, ethoxy group, n-propoxy group, i-propoxy group,

n-butoxy group, 2-methylpropoxy group, 1-methylpropoxy group, and t-butoxy group), a cyano group, a cyanoalkyl group having 2-5 carbon atoms (such as a cyanomethyl group, 2-cyanoethyl group, 3-cyanopropyl group, and 4-cyanobutyl group), an alkylcarbonyloxy group having 2-5 carbon atoms (such as a methylcarbonyloxy group, ethylcarbonyloxy group, and t-butylcarbonyloxy group), an alkoxycarbonyl group having 2-5 carbon atoms (such as a methoxycarbonyl group, ethoxycarbonyl group, and t-butoxycarbonyl group), an alkoxycarbonylalkoxy group having 3-10 carbon atoms (such as a methoxycarbonylmethoxy group, ethoxycarbonylmethoxy group, and t-butoxycarbonylmethoxy group), a halogen atom (such as a fluorine atom and chlorine atom), and a fluoroalkyl group having 1-4 carbon atoms (such as a fluoromethyl group, trifluoromethyl group, and pentafluoroethyl group) can be given. Any number of one or more types of these substituents may be present in Ar¹ or Ar².

[0027]

In the definition of the formula (1), "Ar¹ and Ar² may form, in combination and together with A⁺ in the formula (1), a group having a cyclic structure with 3-8 atoms" refers to a cyclic structure with 3-8 atoms formed from Ar¹ and Ar², wherein two residual groups formed by removing one atom or one group respectively from Ar¹ and Ar² bond directly or via another divalent atom (such as an oxygen atom or a sulfur atom) or another divalent group (for example, an alkylene group such as a methylene group, 1,1-ethylene group, and 1,2-ethylene group or a -NH- group, a -CO- group, or a -SO₂- group) in combination and together with A⁺ in the formula (1).

The number of atoms including A⁺ in this cyclic structure is preferably 5-6. Two or more of the Ar¹ and Ar² groups present in the formulas (1) - (4) may be either the same or different.

[0028]

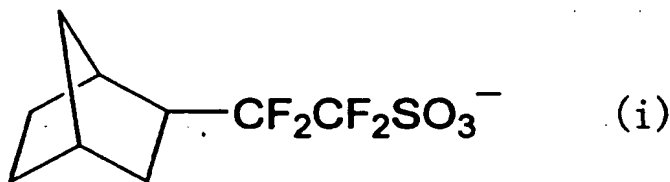
As examples of the anion moiety in the acid generators (A1)-(A4), sulfonic acid anions, MX_a⁻ (wherein M indicates a boron atom, a phosphorus atom, an arsenic atom,

or an antimony atom, X represents a halogen atom, and a is an integer of 4-6), halogen anions, perchloric acid anions, nitric acid anions, phosphoric acid anions, fluoro phosphoric acid anions, and trifluoroacetic acid anions can be given.

Of these anions, the sulfonic acid anions are particularly preferable. Specific examples of the sulfonic acid anions include CH_3SO_3^- , $\text{n-C}_4\text{H}_9\text{SO}_3^-$ (n-butanesulfonic acid anion), $\text{n-C}_8\text{H}_{17}\text{SO}_3^-$ (n-octanesulfonic acid anion), $\text{p-CH}_3\text{C}_6\text{H}_4\text{SO}_3^-$ (p-toluenesulfonic acid anion), CF_3SO_3^- , $\text{n-C}_4\text{F}_9\text{SO}_3^-$ (nonafluoro-n-butanesulfonic acid anion), $\text{n-C}_8\text{H}_{17}\text{SO}_3^-$ (perfluoro-n-octanesulfonic acid anion), and a sulfonic acid anion represented by the following formula (i) (hereinafter referred to as "sulfonic acid anion (i)"), with CF_3SO_3^- , $\text{n-C}_4\text{F}_9\text{SO}_3^-$, and the sulfonic acid anion (i) being particularly preferable.

[0029]

[Chemical Formula 9]



[0030]

Acid generator (A1)

In the various groups represented by R and R' in the formula (1), the number of carbon atoms in the unsubstituted alkyl group having 1-20 carbon atoms is preferably 1-18, more preferably 1-15, and particularly preferably 1-12.

As specific examples of the alkyl group, a methyl group, ethyl group, n-propyl group, i-propyl group, n-butyl group, 2-methylpropyl group, 1-methylpropyl group, t-butyl group, n-pentyl group, n-hexyl group, n-heptyl group, n-octyl group, n-nonyl group, n-decyl group, n-undecyl group, n-dodecyl group, t-dodecyl group, n-tridecyl

group, n-tetradecyl group, n-pentadecyl group, n-hexadecyl group, n-heptadecyl group, n-octadecyl group, n-nonadecyl group; and n-eicosyl group can be given.

[0031]

In the unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon
5 atoms represented by R and R', the carbon atom number is preferably 3-18, more preferably 3-12, and particularly preferably 3-8.

As specific examples of the monovalent alicyclic hydrocarbon group, cycloalkyl groups such as a cyclopropyl group, cyclobutyl group, cyclopentyl group, and cyclohexyl group and bridged alicyclic hydrocarbon groups having 6-20 carbon atoms
10 such as a norbornyl group, tricyclodecanyl group, tetracyclododecyl group, adamantyl group, 3-methyl-1-adamantyl group, 3-methyl-3-adamantyl group, 3-ethyl-1-adamantyl group, 3-ethyl-3-adamantyl group, 3-n-butyl-1-adamantyl group, and 3-n-butyl-3-adamantyl group can be given.

[0032]

15 In the alkenyl group having 2-20 carbon atoms represented by R and R', the carbon atom number is preferably 2-18, more preferably 2-15, and particularly preferably 2-12.

As specific examples of the alkenyl group, a vinyl group, isopropenyl group, 1-propenyl group, and 2-propenyl group can be given.

20 [0033]

In the unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms represented by R and R', the carbon atom number is preferably 6-12, more preferably 6-10, and particularly preferably 6-8.

As specific examples of the monovalent aromatic hydrocarbon group, a phenyl
25 group, o-tolyl group, m-tolyl group, p-tolyl group, benzyl group, o-methylbenzyl group, m-methylbenzyl group, p-methylbenzyl group, 2,3-xylyl group, 2,4-xylyl group, 2,5-xylyl group, 2,6-xylyl group, 3,4-xylyl group, 5,5-xylyl group, mesityl group,

1-naphthyl group, 2-naphthyl group, 1-anthryl group, and 9-anthryl group can be given.

[0034]

The unsubstituted monovalent heterocyclic group having 3-20 atoms represented by R and R' contains one or more hetero atoms such as a nitrogen atom, oxygen atom, and sulfur atom. The number of the atoms in the monovalent heterocyclic group is preferably 4-12, more preferably 5-10, and particularly preferably 5-7.

As specific examples of the monovalent heterocyclic group, 2-pyridyl group, 3-pyridyl group, 4-pyridyl group, 2-imidazolyl group, 4-imidazolyl group, 5-imidazolyl group, 2-morpholinyl group, 3-morpholinyl group, 2-piperidinyl group, 3-piperidinyl group, 4-piperidinyl group, 2-pyrrolidinyl group, and 3-pyrrolidinyl group can be given.

[0035]

As examples of substituents for the substituted alkyl group having 1-20 carbon atoms, substituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, substituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, or substituted monovalent heterocyclic group having 3-20 atoms represented by R or R', the same groups given as examples of the substituents for the substituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms and substituted monovalent heterocyclic group having 3-20 atoms represented by Ar¹ or Ar² can be given. Any number of one or more types of these substituents may be present in R or R'.

[0036]

In the definition of the formula (1), "two R' groups form, in combination and together with the nitrogen atom in the formula, a group having a cyclic structure with 3-8 atoms" relating to the -N(R')₂ group refers to a cyclic structure with 3-8 atoms formed from two R' groups, wherein two residual groups formed by removing one atom or one group from the two R' groups bond directly or via another divalent atom (such as an oxygen atom or a sulfur atom) or another divalent group (for example, an alkylene group such as a methylene group, 1,2-ethylene group, or a -NH- group, a -CO- group, or

a -SO₂- group) in combination and together with the nitrogen atom in the formula.

The number of atoms including the nitrogen atom in this cyclic structure is preferably 5-6.

[0037]

- 5 As specific examples of preferable R or R' groups, a methyl group, ethyl group, n-propyl group, i-propyl group, n-butyl group, t-butyl group, n-pentyl group, n-hexyl group, n-heptyl group, n-octyl group, n-dodecyl group, t-dodecyl group, n-hexadodecyl group, cyclopentyl group, cyclohexyl group, norbornyl group, p-tolyl group, benzyl group, phenyl group, 1-naphthyl group, 2-naphthyl group, trifluoromethyl group, nonafluoro-n-butyl group, perfluoro-n-octyl group, methoxycarbonyldifluoromethyl group, and campholoyl group can be given.
- 10

The group -N(R')₂, particularly a group such as -N(CH₃)₂ and -N(C₂H₅)₂, is also a preferable R group.

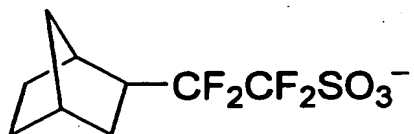
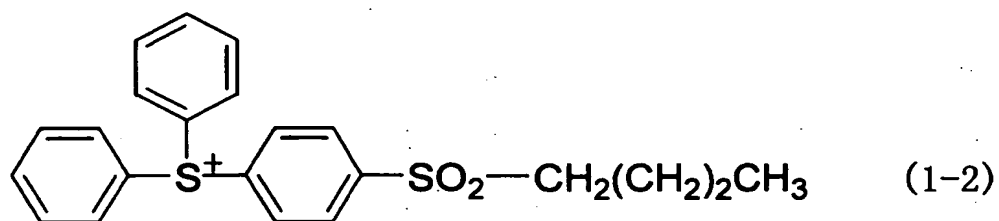
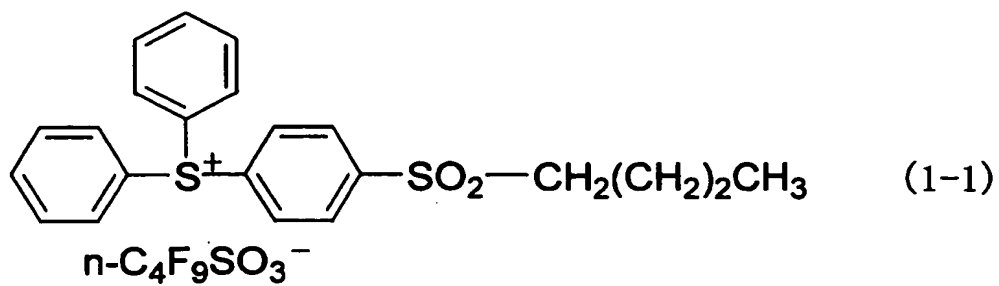
- When two or more R groups are present in the compound of formula (1), such groups may be either identical or different.
- 15

[0038]

The compounds of the following formulas (1-1) to (1-14) can be given as specific preferable examples of the onium salt compound (1).

[0039]

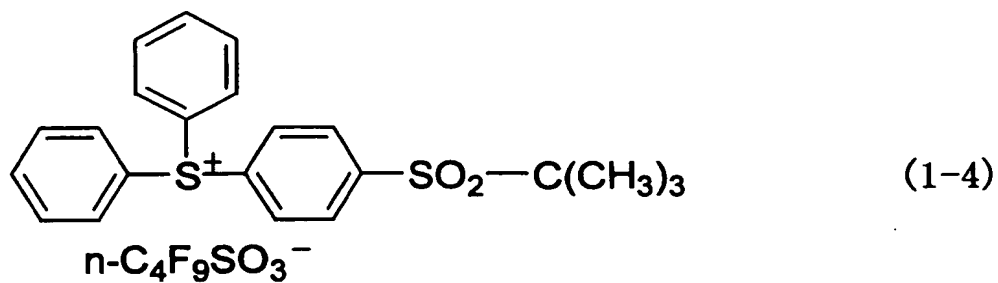
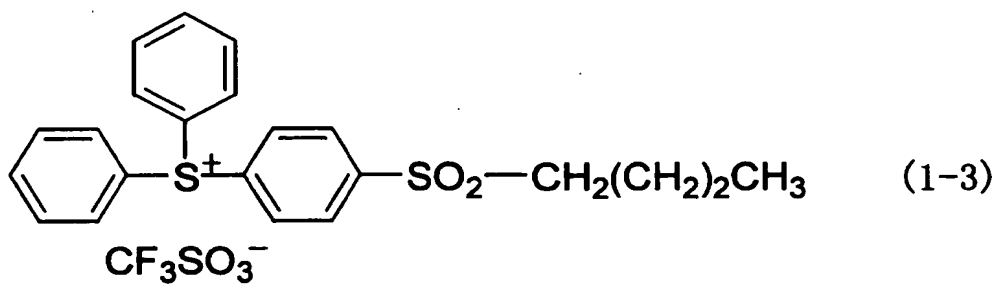
- 20 [Chemical Formula 10]



[0040]

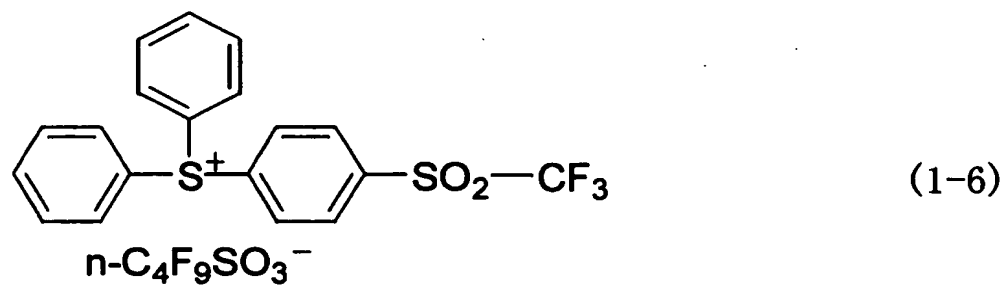
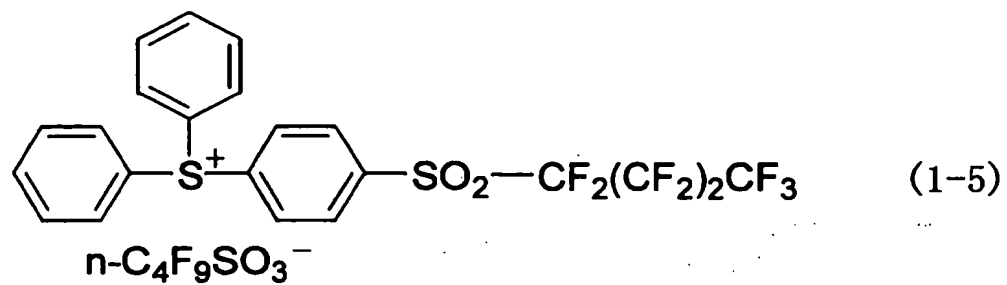
[Chemical Formula 11]

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[0041]

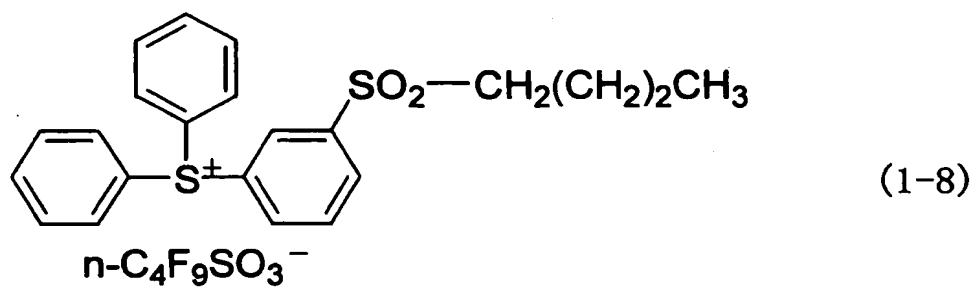
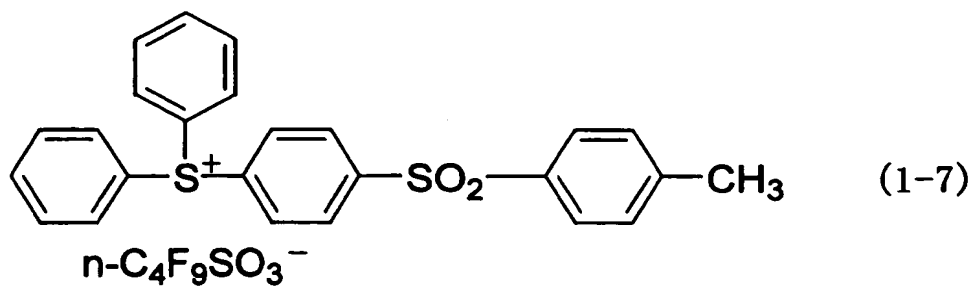
[Chemical Formula 12]



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[0042]

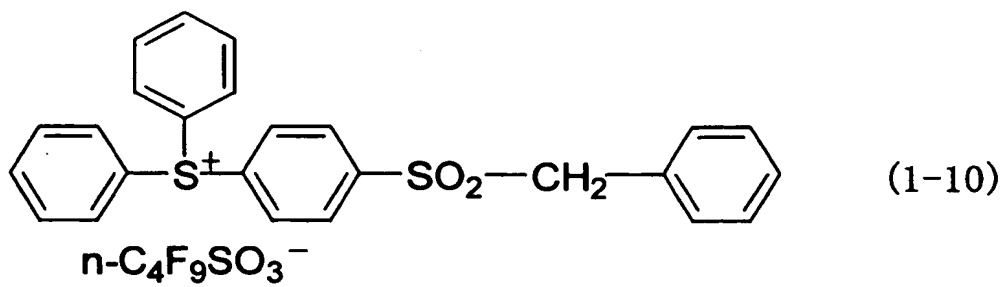
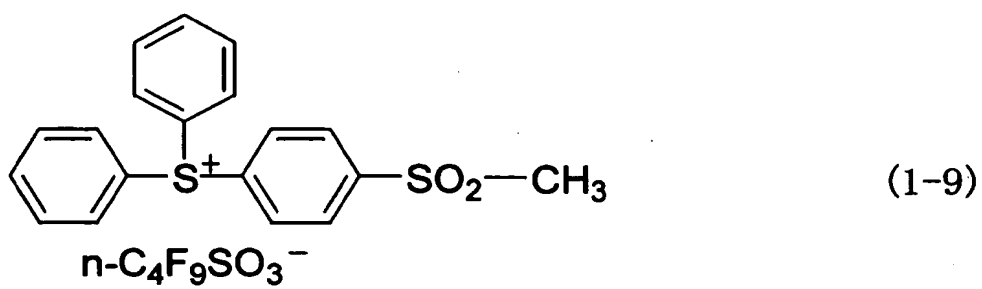
[Chemical Formula 13]



[0043]

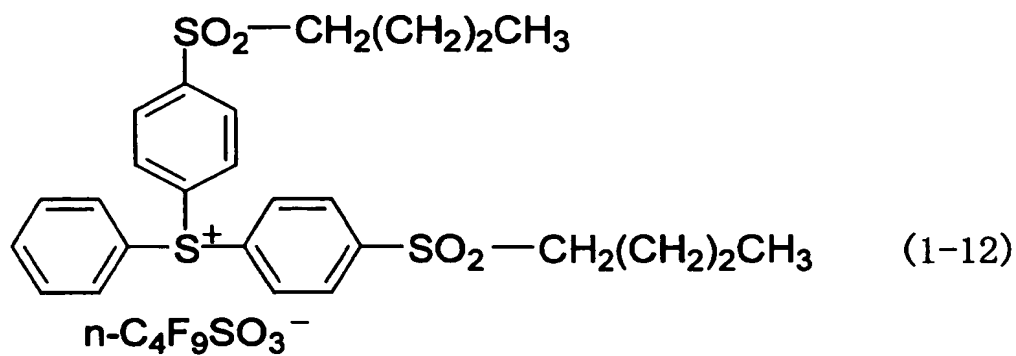
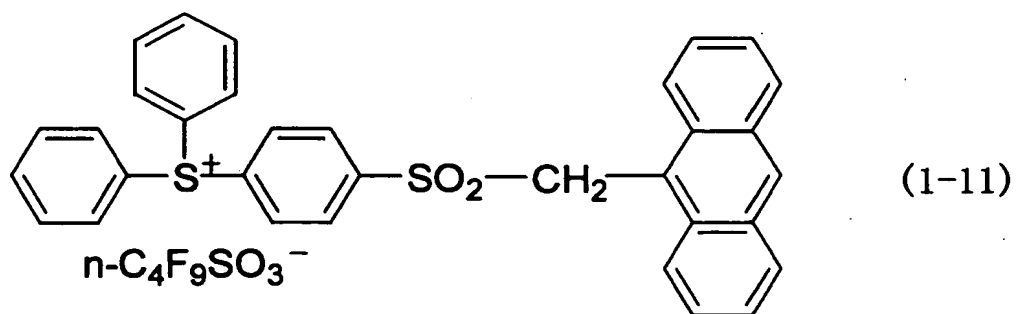
[Chemical Formula 14]

5



[0044]

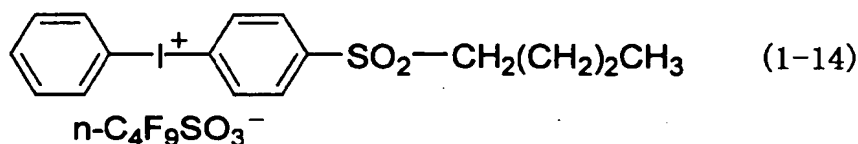
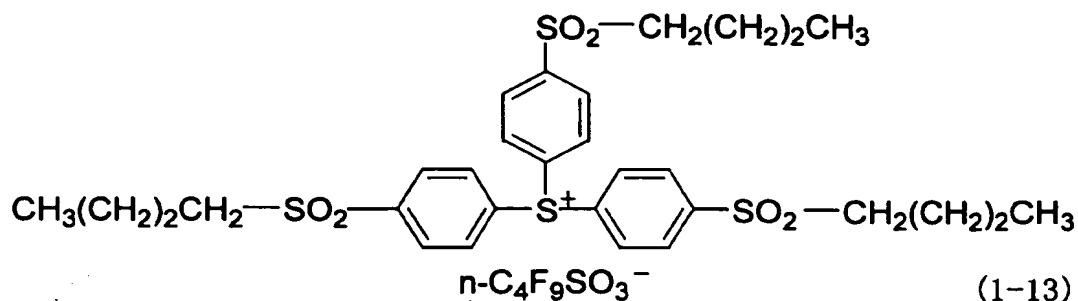
[Chemical Formula 15]



5

[0045]

[Chemical Formula 16]



[0046]

The acid generator (A1) is suitable for use as a photoacid generator responsive to active radiations such as (deep) ultraviolet rays represented by a KrF excimer laser, ArF excimer laser, F₂ excimer laser, and EUV, as well as to electron beams, in a chemically-amplified photoresist used in the field of microfabrication represented by the manufacture of integrated circuit devices.

[0047]

10 Onium salt compound (2)

As examples of the unsubstituted alkyl group having 1-20 carbon atoms, unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, alkenyl group having 2-20 carbon atoms, unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, and unsubstituted monovalent heterocyclic group having 3-20 atoms represented by R¹ and R'' in the formula (2), the same groups as given for the corresponding groups for R and R' in the formula (1) can be given.

As examples of the substituents for these groups, the same groups as previously mentioned in connection with the substituted monovalent aromatic hydrocarbon groups

having 6-20 carbon atoms and substituted monovalent heterocyclic groups having 3-20 atoms represented by Ar^1 and Ar^2 can be given. Any number of one or more types of these substituents may be present in R^1 or R'' .

[0048]

5 In the definition of R^1 , "two R'' groups form, in combination and together with the nitrogen atom in the formula, a group having a cyclic structure with 3-8 atoms" relating to the $-N(R'')_2$ group refers to a cyclic structure with 3-8 atoms formed from two R'' groups, wherein two residual groups formed by removing one atom or one group from the two R'' groups bond directly or via another divalent atom (such as a
10 nitrogen atom, oxygen atom, or sulfur atom) or another divalent group (for example, an alkylene group such as a methylene group, 1,2-ethylene group, or a $-NH-$ group, a $-CO-$ group, or a $-SO_2-$ group) in combination and together with the nitrogen atom in the formula.

The number of atoms including the nitrogen atom in this cyclic structure is
15 preferably 5-6.

[0049]

The same preferable specific examples given for R in the formula (1) can be given as preferable specific examples for R^1 in the formula (2), and the same preferable specific examples given for R' in formula (1) can be given as preferable specific
20 examples for R'' .

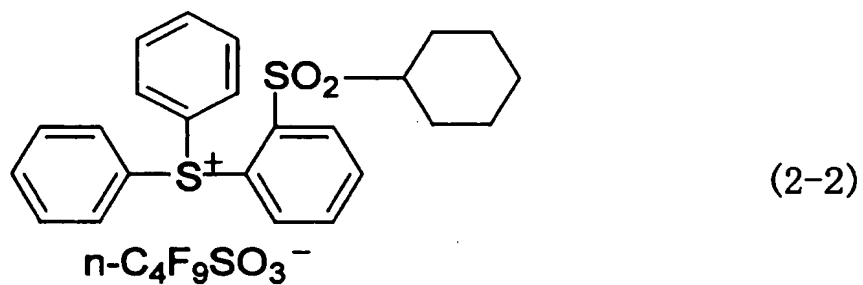
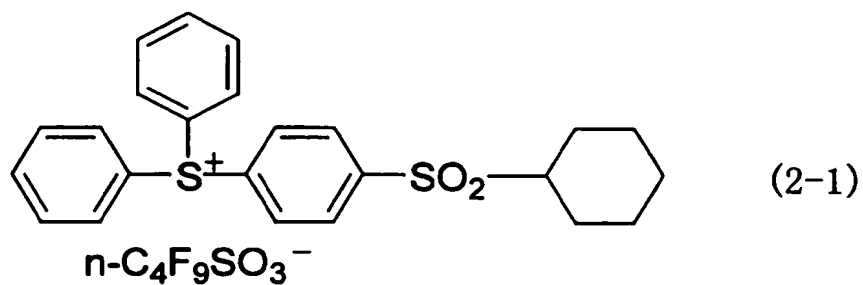
When two or more R^1 groups are present in the compound of formula (2), such groups may be either identical or different. In the formula (2), p is preferably 0-2 and q is preferably 1-3.

[0050]

25 The compounds of the following formulas (2-1) to (2-4) can be given as specific preferable examples of the onium salt compound (2).

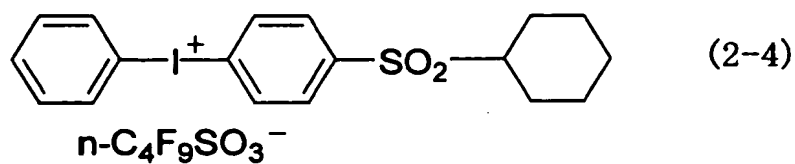
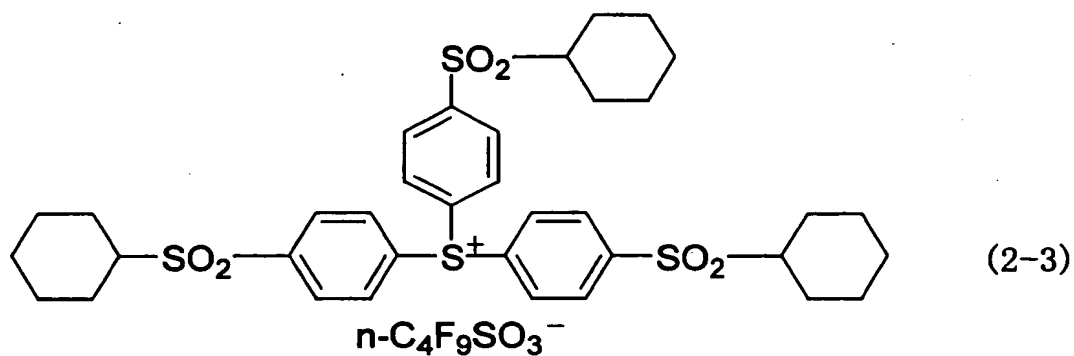
[0051]

[Chemical Formula 17]



5 [0052]

[Chemical Formula 18]



[0053]

The onium salt compound (2) is not only extremely suitable for use as a photoacid generator responsive to active radiations such as (deep) ultraviolet rays represented by a KrF excimer laser, ArF excimer laser, F₂ excimer laser, and EUV, as well as to electron beams, in a chemically-amplified photoresist used in microfabrication represented by the manufacture of integrated circuit devices, but also useful as a raw material for the synthesis of a heat acid generator which generates an acid with heating and other related onium salt compounds.

[0054]

Onium salt compound (3)

As examples of the unsubstituted alkyl group having 1-20 carbon atoms, unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms, alkenyl group having 2-20 carbon atoms, unsubstituted monovalent aromatic hydrocarbon group having 6-20 carbon atoms, and unsubstituted monovalent heterocyclic group having 3-20 atoms represented by R² and R''' in the formula (3), the same groups as given for the corresponding groups for R and R' in the formula (1) can be given.

As examples of the substituents for these groups, the same groups as previously mentioned in connection with the substituted monovalent aromatic hydrocarbon groups having 6-20 carbon atoms and substituted monovalent heterocyclic groups having 3-20 atoms represented by Ar¹ and Ar² can be given. Any number of one or more types of these substituents may be present in R² or R'''.

[0055]

In the definition of R², "two R'" groups form, in combination and together with the nitrogen atom in the formula, a group having a cyclic structure with 3-8 atoms" relating to the -N(R''')₂ group refers to a cyclic structure with 3-8 atoms formed from

two R''' groups, wherein two residual groups formed by removing one atom or one group from the two R''' groups bond directly or via another divalent atom (such as a nitrogen atom, oxygen atom, or sulfur atom) or another divalent group (for example, an alkylene group such as a methylene group, 1,2-ethylene group, or a -NH- group, a -CO- group, or a -SO₂- group) in combination and together with the nitrogen atom in the formula.

The number of atoms including the nitrogen atom in this cyclic structure is preferably 5-6.

[0056]

10 The same preferable specific examples given for R in the formula (1) can be given as preferable specific examples for R² in the formula (3), and the same preferable specific examples given for R' in formula (1) can be given as preferable specific examples for R'''.

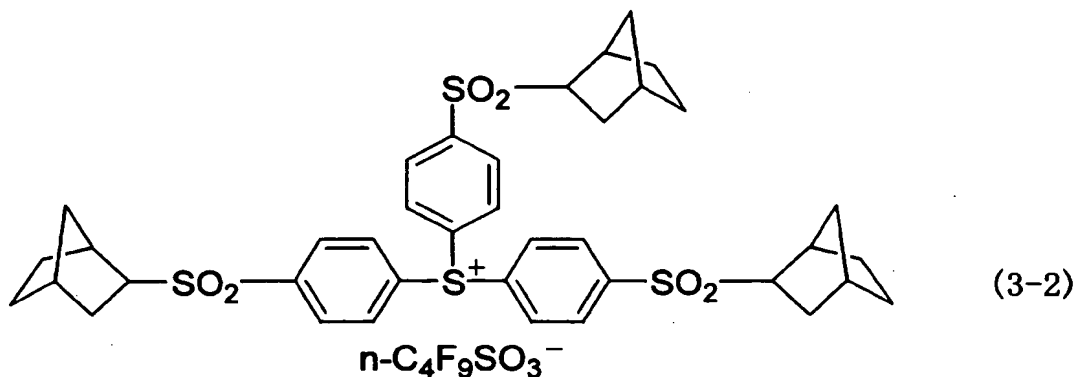
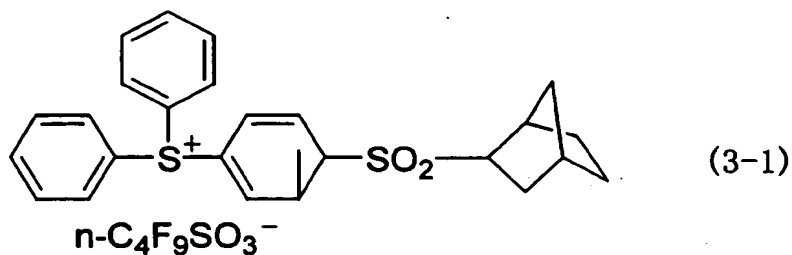
15 When two or more R² groups are present in the compound of formula (3), such groups may be either identical or different. In the formula (3), r is preferably 0-2, s is preferably 0-2, and v is preferably 1 or 2.

[0057]

The compounds of the following formulas (3-1) to (3-2) can be given as specific preferable examples of the onium salt compound (3).

20 [0058]

[Chemical Formula 19]



[0059]

The onium salt compound (3) is not only extremely suitable for use as a
 5 photoacid generator responsive to active radiations such as (deep) ultraviolet rays
 represented by a KrF excimer laser, ArF excimer laser, F₂ excimer laser, and EUV, as
 well as to electron beams, in a chemically-amplified photoresist used in
 microfabrication represented by the manufacture of integrated circuit devices, but also
 useful as a raw material for the synthesis of a heat acid generator which generates an
 10 acid with heating and other related onium salt compounds.

[0060]

Onium salt compound (4)

As examples of the unsubstituted alkyl group having 1-20 carbon atoms and
 unsubstituted monovalent alicyclic hydrocarbon group having 3-20 carbon atoms
 15 represented by R³ or R⁴ in the formula (4), the same groups as given for the
 corresponding groups for R and R' in the formula (1) can be given.

As examples of the substituents for these groups, the same groups as previously mentioned in connection with the substituted aryl groups having 6-20 carbon atoms and substituted monovalent heterocyclic groups having 3-20 atoms represented by Ar¹ and Ar² can be given. Any number of one or more types of these substituents may be present in R³ or R⁴.

[0061]

In the definition of the formula (4), "R³ and R⁴ may form, in combination and together with one carbon atom and two oxygen atoms in the formula, a group having a cyclic structure with 4-10 atoms" refers to a cyclic structure with 4-10 atoms formed from the R³ and R⁴ groups, wherein two residual groups formed by removing one atom or one group from each of the R³ group and R⁴ group bond directly or via another divalent atom (such as an oxygen atom or a sulfur atom) or another divalent group (for example, an alkylene group such as a methylene group, 1,2-ethylene group, 1,2-dimethyl-1,2-ethylene group, trimethylene group, or a -NH- group, a -CO- group, or a -SO₂- group) in combination and together with one carbon atom and two oxygen atoms in the formula.

The number of atoms in this cyclic structure including the one carbon atom and two oxygen atoms is preferably 5-6.

[0062]

As specific examples of preferable R³ or R⁴ groups in the formula (4), a methyl group, ethyl group, n-propyl group, i-propyl group, n-butyl group, t-butyl group, n-pentyl group, n-hexyl group, n-heptyl group, n-octyl group, n-dodecyl group, t-dodecyl group, n-hexadodecyl group, cyclopentyl group, cyclohexyl group, norbornyl group, p-tolyl group, benzyl group, phenyl group, 1-naphthyl group, 2-naphthyl group, trifluoromethyl group, nonafluoro-n-butyl group, perfluoro-n-octyl group, methoxycarbonyl difluoromethyl group, and campholoyl group, as well as groups formed by bonding of the R³ group and R⁴ group, such as 1,2-ethylene group,

1,2-dimethyl-1,2-ethylene group, trimethylene group, 3,4-tetrahydrofuranylene group, and 3,4-tetrahydropyranylene group can be given.

When two or more R^3 groups and R^4 groups are present in the compound of formula (4), such groups may be either identical or different.

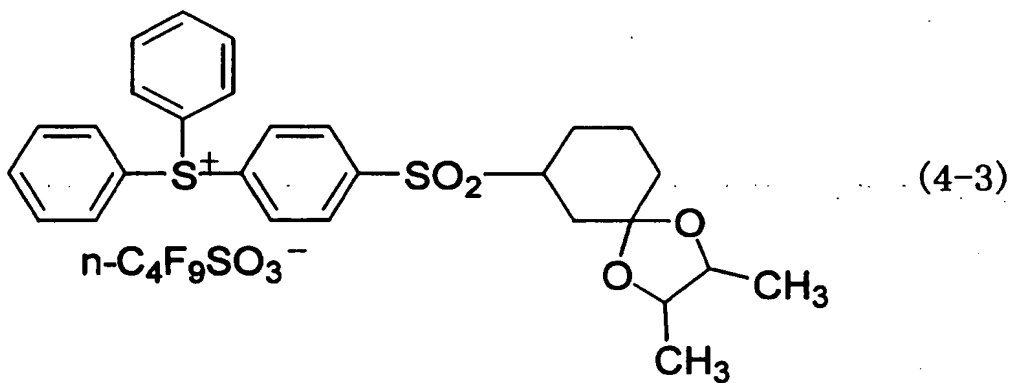
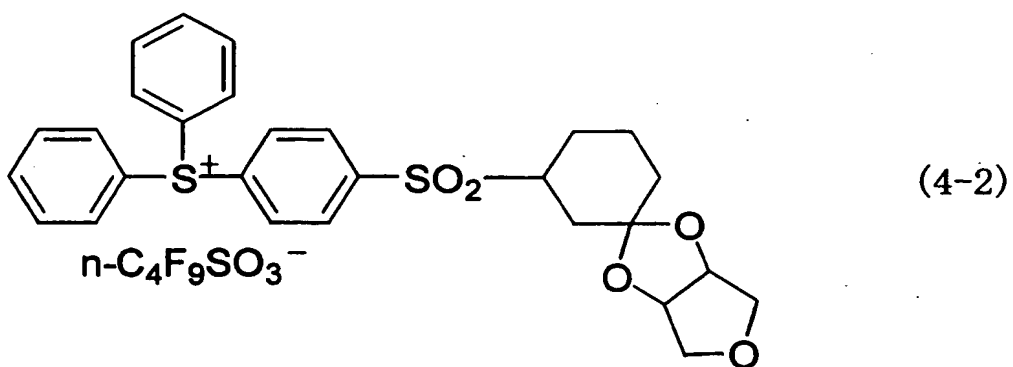
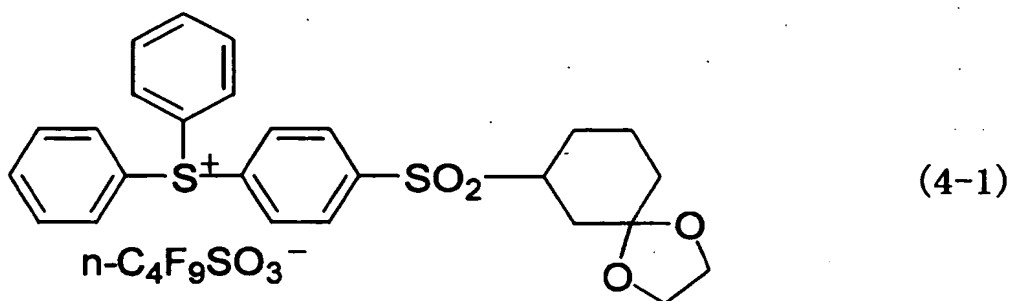
5 In the formula (4), t and s are preferably 1-3, and more preferably 3.

[0063]

The compounds of the following formulas (4-1) to (4-7) can be given as specific preferable examples of the onium salt compound (4).

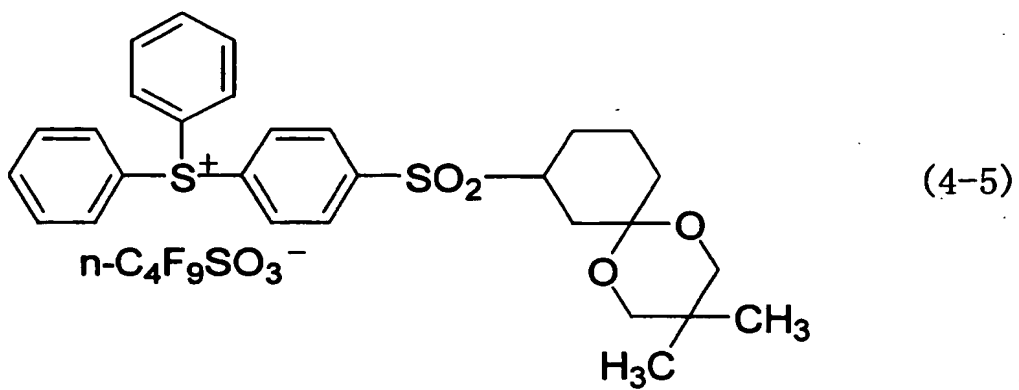
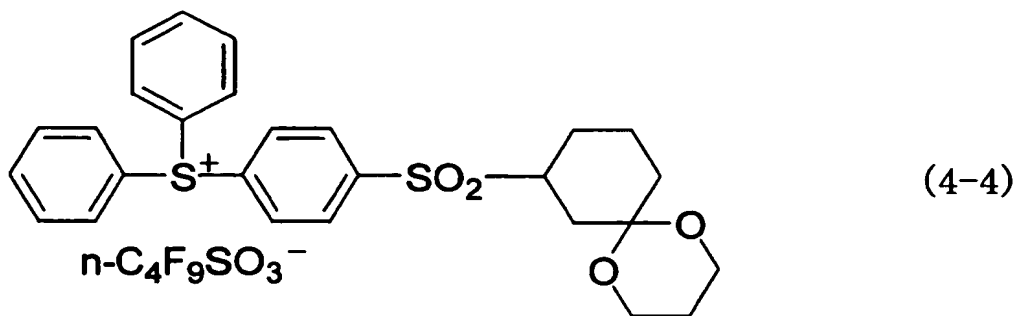
[0064]

10 [Chemical Formula 20]



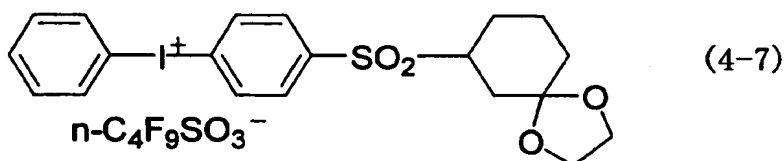
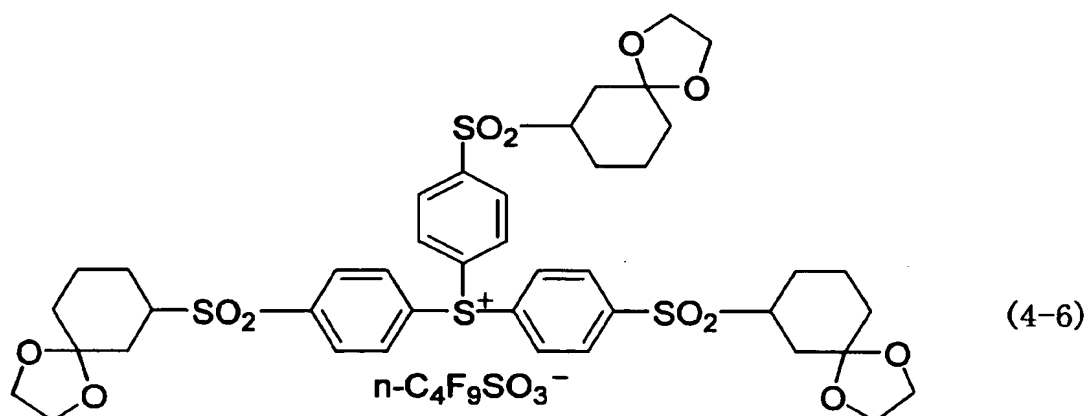
[0065]

[Chemical Formula 21]



[0066]

[Chemical Formula 22]



[0067]

The onium salt compound (4) is not only extremely suitable for use as a
 5 photoacid generator responsive to active radiations such as (deep) ultraviolet rays
 represented by a KrF excimer laser, ArF excimer laser, F₂ excimer laser, and EUV, as
 well as to electron beams, in a chemically-amplified photoresist used in
 microfabrication represented by the manufacture of integrated circuit devices, but also
 useful as a raw material for the synthesis of a heat acid generator which generates an
 10 acid with heating and other related onium salt compounds.

[0068]

Synthesis of onium salt compounds (1)-(4)

The onium salt compound (1) can be synthesized by the reaction formula shown
 below according to the method described in non-patent document 2, for example.

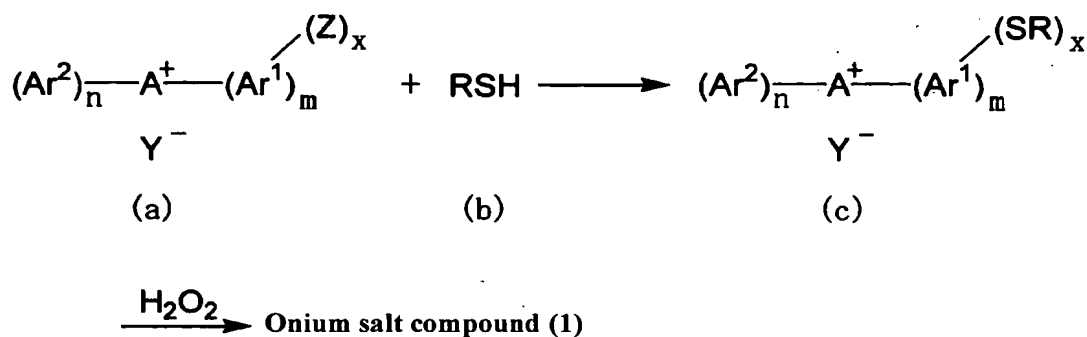
15 Specifically, the onium salt compound precursor (a) is reacted with a thiol
 compound (b) in the presence of a basic compound (hereinafter referred to as “a
 thioetherification reaction”) to convert the onium salt compound precursor (a) into an

onium salt compound (c) having an -SR group, which is oxidized by an oxidizing agent such as hydrogen peroxide solution to produce the onium salt compound (1). The onium salt compounds (2)-(4) can also be synthesized in the same manner as in the synthesis of the onium salt compound (1).

5 [0069]

[Chemical Formula 23]

Reaction Formula



[0070]

10 wherein Z is a dissociable monovalent group bonded to the Ar¹ group, A, Ar¹, m, Ar², n, R, and x are respectively the same as A, Ar¹, m, Ar², n, R, and x in the formula (1), and Y⁻ represents a monovalent anion.

[0071]

[Non-patent document 2]

15 J. Org. Chem., Vol. 48, p. 605-609 (1983)

[0072]

As examples of the dissociable monovalent group represented by Z of the onium salt compound precursor (a), in addition to halogen atoms such as fluorine, chlorine, bromine, and iodine, a CH₃SO₃⁻ group, a p-CH₃C₆H₄SO₃⁻ group (a p-toluenesulfonate residue), and the like can be given, with fluorine and chlorine atoms being preferable.

20

[0073]

The molar ratio of the thiol compound (b) to the onium salt compound precursor (a) during the thioetherification reaction is usually 1-100 and preferably 1.5-10.

[0074]

5 As examples of the basic compound used in the thioetherification reaction, potassium hydroxide, sodium hydroxide, lithium carbonate, sodium carbonate, potassium carbonate, lithium hydrogencarbonate, sodium hydrogencarbonate, potassium hydrogencarbonate, and the like can be given. Of these, sodium hydroxide and potassium hydroxide are preferable.

10 The molar ratio of the basic compound to the thiol compound (b) in the thioetherification reaction is usually 1.0-10.0 and preferably 2.0-4.0.

[0075]

The thioetherification reaction is preferably carried out in a mixed solvent of an organic solvent and water. As the organic solvent, dichloromethane, chloroform, 15 toluene, benzene, hexane, ethyl acetate, and the like are preferable. Of these, dichloromethane and chloroform are particularly preferable.

The amount of the organic solvent used is usually 5 parts by weight or more, preferably 10 parts by weight or more, and more preferably 20-90 parts by weight for 100 parts by weight of the total amount of the organic solvent and water.

20 [0076]

The thioetherification reaction is carried out at a temperature of usually -40°C to +50°C, and preferably -20°C to +30°C, for usually 0.1-72 hours, and preferably 0.5-3 hours. If the reaction temperature used is higher than the boiling point of the organic solvent or water, a pressure vessel such as an autoclave is used.

25 [0077]

As the oxidizing agent used in the oxidation reaction of the onium salt compound (c), in addition to hydrogen peroxide, methachloroperbenzoic acid, t-butyl

hydroperoxide, potassium peroxysulfate, potassium permanganate, sodium perborate, sodium metaiodate, chromic acid, sodium dichromate, halogen, iodobenzene dichloride, iodobenzene diacetate, osmium oxide (VII), ruthenium oxide (VII), sodium hypochlorite, sodium chlorite, oxygen gas, ozone gas, and the like can be given, with
5 hydrogen peroxide, methachloroperbenzoic acid, and t-butyl hydroperoxide being preferable.

The molar ratio of the oxidizing agent to the onium salt compound (c) is usually 1.0-20.0 and preferably 1.5-10.0.

[0078]

10 A transition metal catalyst may be used together with the oxidizing agent in the oxidation reaction.

As examples of the transition metal catalyst, disodium tungstate, iron (III) chloride, ruthenium (III) chloride, and selenium (IV) oxide can be given, with disodium tungstate being preferable.

15 The molar ratio of the transition metal catalyst to the onium salt compound (7) is usually 0.001-2.0, preferably 0.01-1.0, and particularly preferably 0.03-0.5.

[0079]

Furthermore, in addition to the oxidizing agent and the transition metal catalyst, a buffer agent may be used in the oxidation reaction to control the pH of the reaction
20 solution.

As examples of the buffer agent, disodium hydrogenphosphate, sodium dihydrogenphosphate, dipotassium hydrogenphosphate, potassium dihydrogenphosphate, and the like can be given. The molar ratio of the buffer agent to the onium salt compound (7) is usually 0.01-2.0, preferably 0.03-1.0, and particularly
25 preferably 0.05-0.5.

[0080]

The oxidation reaction is usually carried out in a solvent.

As the solvent, water, organic solvents such as lower alcohols, tetrahydrofuran, N,N-dimethylformamide, N,N-dimethylacetamide, acetonitrile, dimethyl sulfoxide, acetic acid, trifluoroacetic acid, and the like can be given as preferable examples, with methanol, N,N-dimethylacetamide, acetonitrile, and dimethyl sulfoxide being more
5 preferable, and methanol being particularly preferable.

The amount of the solvent used is usually 5-100 parts by weight, preferably 10-100 parts by weight, and particularly preferably 20-50 parts by weight for 100 parts by weight of the onium salt compound (c).

If necessary, the organic solvent may be used with water. In this case, the
10 amount of the organic solvent used is usually 5 parts by weight or more, preferably 10 parts by weight or more, and particularly preferably 20-90 parts by weight, for 100 parts by weight of the total amount of the organic solvent and water.

[0081]

The oxidation reaction is carried out at a temperature of usually 0-100°C,
15 preferably 5-60°C, and more preferably 5-40°C, for usually 0.1-72 hours, and preferably 0.5-24 hours.

[0082]

Positive-tone radiation-sensitive resin composition

Acid generator (A)

20 The component (A) of the positive-tone radiation-sensitive resin composition of the present invention is a photoacid generator (hereinafter referred to as "acid generator (A)") comprising at least one type of photoacid generator as an essential component, selected from the group consisting of acid generators (A1)-(A4).

In the acid generators (A1)-(A4), A, Ar¹, m, Ar², n, and x in the formulas (1)-(4)
25 are individually the same or different. The acid generators (A1)-(A4) can be used either individually or in combination of two or more in the positive-tone radiation-sensitive resin composition of the present invention.

[0083]

One or more photoacid generators other than the acid generators (A1)-(A4) (hereinafter referred to as "other acid generators") can be used in combination in the positive-tone radiation-sensitive resin composition of the present invention.

5 As examples of the other acid generators, onium salt compounds, sulfone compounds, sulfonate compounds, sulfonimide compounds, diazomethane compounds, disulfonylmethane compounds, and the like can be given.

[0084]

10 As examples of the onium salt compounds, iodonium salts, sulfonium salts (including tetrahydrothiophenium salts), phosphonium salts, diazonium salts, ammonium salts, pyridinium salts, and the like can be given.

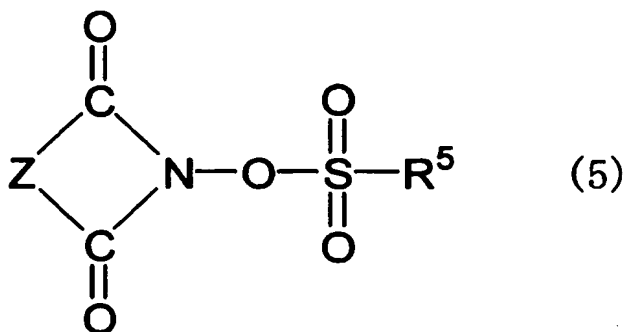
As examples of the sulfone compound, β -ketosulfone, β -sulfonylsulfone, and α -diazo compounds of these compounds can be given.

15 As examples of the sulfonate compound, alkyl sulfonate, haloalkyl sulfonate, aryl sulfonate, and imino sulfonate can be given.

As an example of the sulfonimide compound, a compound of the following formula (5), and the like can be given,

[0085]

[Chemical Formula 24]



20 wherein Z is a divalent organic group and R^5 is a monovalent organic group.

[0086]

As examples of z in the formula (5), a methylene group, linear or branched alkylene group having 2-20 carbon atoms, aralkylene group having 2-20 carbon atoms, difluoromethylene group, linear or branched perfluoroalkylene group having 2-20 carbon atoms, cyclohexylene group, phenylene group, substituted or unsubstituted divalent group possessing a norbornene skeleton, or a group wherein these groups are substituted with an aryl group having six or more carbon atoms or an alkoxyl group having one or more carbon atoms can be given.

[0087]

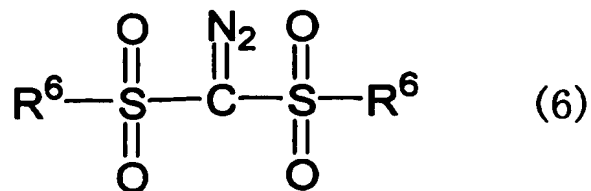
As examples of R⁵, a linear or branched alkyl group having 1-10 carbon atoms, linear or branched perfluoroalkyl group having 1-10 carbon atoms, perfluorocycloalkyl group having 3-10 carbon atoms, monovalent hydrocarbon group possessing a bicyclo ring having 7-15 carbon atoms, and an aryl group having 6-12 carbon atoms, can be given.

[0088]

As an example of the diazomethane compound, a compound of the following formula (6) can be given,

[0089]

[Chemical Formula 25]



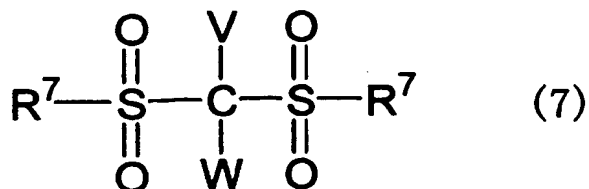
wherein R⁶ individually represents a monovalent group such as a linear or branched alkyl group, cycloalkyl group, aryl group, halogenated alkyl group, halogenated cycloalkyl group, and halogenated aryl group.

[0090]

As an example of the disulfonylmethane compound, a compound of the following formula (7) can be given,

[0091]

5 [Chemical Formula 26]



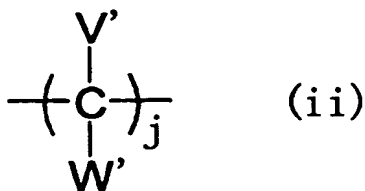
[0092]

- 10 wherein R⁷ individually represents a linear or branched monovalent aliphatic hydrocarbon group, cycloalkyl group, aryl group, aralkyl group, or other monovalent organic group having a hetero atom, V and W individually represent an aryl group, hydrogen atom, linear or branched monovalent aliphatic hydrocarbon group, cycloalkyl group, aralkyl group, or other monovalent organic group having a hetero atom, provided
- 15 that at least one of V and W represents an aryl group, or V and W bond to form a monocyclic or polycyclic ring having at least one unsaturated bond, or V and W bond to form a group shown by the following formula (ii),

[0093]

[Chemical Formula 27]

20



[0094]

wherein V's and W's individually represent a hydrogen atom, halogen atom, linear or branched alkyl group, cycloalkyl group, aryl group, or aralkyl group, or V' and W', each
 5 bonded to the same or different carbon atoms, may bond to form a carbomonocyclic structure, two or more of the V' and W', if present, may be either the same or different, and j is an integer from 2 to 10.

[0095]

The other acid generator is preferably one or more acid generators selected from
 10 the group consisting of an onium salt compound, sulfonimide compound, and diazomethane compound.

As a particularly preferable other acid generator, at least one compound selected from the group consisting of bis(4-t-butylphenyl)iodonium trifluoromethanesulfonate, bis(4-t-butylphenyl)iodonium nonafluoro-n-butanesulfonate,
 15 bis(4-t-butylphenyl)iodonium p-toluenesulfonate, bis(4-t-butylphenyl)iodonium 10-camphorsulfonate, bis(4-t-butylphenyl)iodonium 2-trifluoromethylbenzenesulfonate, bis(4-t-butylphenyl)iodonium 4-trifluoromethylbenzenesulfonate, bis(4-t-butylphenyl)iodonium 2,4-difluorobenzenesulfonate, triphenylsulfonium trifluoromethanesulfonate, triphenylsulfonium nonafluoro-n-butanesulfonate,
 20 triphenylsulfonium p-toluenesulfonate, triphenylsulfonium 10-camphorsulfonate, triphenylsulfonium 2-trifluoromethylbenzenesulfonate, triphenylsulfonium 4-trifluorobenzenesulfonate, triphenylsulfonium 2,4-difluoromethylbenzenesulfonate, 1-(4-n-butoxynaphthalene-1-yl)tetrahydrothiophenium trifluoromethanesulfonate, 1-(4-n-butoxynaphthalene-1-yl)tetrahydrothiophenium nonafluoro-n-butanesulfonate,
 25 N-(trifluoromethanesulfonyloxy)succinimide, N-(trifluoromethanesulfonyloxy)bicyclo[2.2.1]hept-5-ene-2,3-dicarboxyimide, N-(10-camphorsulfonyloxy)succinimide,

N-(10-camphorsulfonyloxy)bicyclo[2.2.1]hept-5-ene-2,3-dicarboxyimide,
N-{(5-methyl-5-carboxymethanebicyclo[2.2.1]hept-2-yl)sulfonyloxy}succinimide,
bis(cyclohexanesulfonyl)diazomethane, bis(t-butylsulfonyl)diazomethane, and
bis(1,4-dioxaspiro[4.5]-decan-7-sulfonyl)diazomethane can be given.

5 [0096]

The proportion of other acid generators can be appropriately determined depending on the types of each other acid generator. The proportion is usually 95 parts by weight or less, preferably 90 parts by weight or less, and particularly preferably 80 parts by weight or less for 100 parts by weight of the total amount of the acid generators (A1) - (A4) and the other acid generators. If the proportion of the other acid generators exceeds 95 parts by weight, the desired effects of the present invention may be impaired.

[0097]

Acid-dissociable group-containing resin (B)

15 The component (B) of the positive-tone radiation-sensitive resin composition of the present invention is an acid-dissociable group-containing resin which is insoluble or scarcely soluble in alkali, but becomes easily soluble in alkali when the acid-dissociable group dissociates (hereinafter referred to as "acid-dissociable group-containing resin (B)").

20 If 50% or more of the initial film thickness of a resist film remains after development when a resist film made only from the acid-dissociable group-containing resin (B) is developed under the same alkaline development conditions employed for forming a resist pattern using a resist film formed from a radiation-sensitive resin composition comprising the acid-dissociable group-containing resin (B), such a
25 characteristic of the acid-dissociable group-containing resin (B) is referred to as "insoluble or scarcely soluble in alkali" in the present invention.

[0098]

The acid-dissociable group of the acid-dissociable group-containing resin (B) refers to a group which is replaced the hydrogen atom in an acid-functional group such as a phenolic hydroxyl group, carboxyl group, and sulfonic group and is dissociable in the presence of an acid.

5 As examples of such an acid-dissociable group, a substituted methyl group, 1-substituted ethyl group, 1-substituted n-propyl group, 1-branched alkyl group, alkoxy carbonyl group, acyl group, cyclic acid-dissociable group, and the like can be given.

[0099]

10 As examples of the substituted methyl group, a methoxymethyl group, methylthiomethyl group, ethoxymethyl group, ethylthiomethyl group, methoxyethoxymethyl group, benzyloxymethyl group, benzylthiomethyl group, phenacyl group, 4-bromophenacyl group, 4-methoxyphenacyl group, 4-methylthiophenacyl group, α -methylphenacyl group, cyclopropylmethyl group, 15 benzyl group, diphenylmethyl group, triphenylmethyl group, 4-bromobenzyl group, 4-nitrobenzyl group, 4-methoxybenzyl group, 4-methylthiobenzyl group, 4-ethoxybenzyl group, 4-ethylthiobenzyl group, piperonyl group, methoxycarbonylmethyl group, ethoxycarbonylmethyl group, n-propoxycarbonylmethyl group, i-propoxycarbonylmethyl group, n-butoxycarbonylmethyl group, and 20 t-butoxycarbonylmethyl group can be given.

[0100]

As examples of the 1-substituted ethyl group, a 1-methoxyethyl group, 1-methylthioethyl group, 1,1-dimethoxyethyl group, 1-ethoxyethyl group, 1-ethylthioethyl group, 1,1-diethoxyethyl group, 1-phenoxyethyl group, 25 1-phenylthioethyl group, 1,1-diphenoxyethyl group, 1-benzyloxyethyl group, 1-benzylthioethyl group, 1-cyclopropyloxyethyl group, 1-cyclohexyloxyethyl group, 1-phenylethyl group, 1,1-diphenylethyl group, 1-methoxycarbonylethyl group,

1-ethoxycarbonylethyl group, 1-n-propoxycarbonylethyl group,
1-i-propoxycarbonylethyl group, 1-n-butoxycarbonylethyl group, and
1-t-butoxycarbonylethyl group can be given.

[0101]

5 As examples of the 1-substituted n-propyl group, a 1-methoxy-n-propyl group
and 1-ethoxy-n-propyl group can be given.

As examples of the 1-branched alkyl group, an i-propyl group, 1-methylpropyl
group, t-butyl group, 1,1-dimethylpropyl group, 1-methylbutyl group, and
1,1-dimethylbutyl group can be given.

10 [0102]

As examples of the alkoxycarbonyl group, a methoxycarbonyl group,
ethoxycarbonyl group, i-propoxycarbonyl group, t-butoxycarbonyl group, and the like
can be given.

[0103]

15 As examples of the acyl group, an acetyl group, propionyl group, butyryl group,
heptanoyl group, hexanoyl group, valeryl group, pivaloyl group, isovaleryl group,
lauryloyl group, myristoyl group, palmitoyl group, stearoyl group, oxalyl group,
malonyl group, succinyl group, glutaryl group, adipoyl group, piperoyl group, suberoyl
group, azelaoyl group, sebacoyl group, acryloyl group, propioloyl group, methacryloyl
20 group, crotonoyl group, oleoyl group, maleoyl group, fumaroyl group, mesaconoyl
group, camphoroyl group, benzoyl group, phthaloyl group, isophthaloyl group,
terephthaloyl group, naphthoyl group, toluoyl group, hydroatropoyl group, atropoyl
group, cinnamoyl group, furoyl group, thenoyl group, nicotinoyl group, isonicotinoyl
group, p-toluenesulfonyl group, and mesyl group can be given.

25 [0104]

As examples of the cyclic acid-dissociable group, a cyclopropyl group,
cyclopentyl group, cyclohexyl group, cyclohexenyl group, 4-methoxycyclohexyl group,

tetrahydropyranyl group, tetrahydrofuranyl group, tetrahydrothiopyranyl group, tetrahydrothiofuranyl group, 3-bromotetrahydropyranyl group, 4-methoxytetrahydropyranyl group, 4-methoxytetrahydrothiopyranyl group, and 3-tetrahydrothiophene-1,1-dioxide group can be given.

5 [0105]

Of these acid-dissociable groups, a benzyl group, t-butoxycarbonylmethyl group, 1-methoxyethyl group, 1-ethoxyethyl group, 1-cyclohexyloxyethyl group, 1-ethoxy-n-propyl group, t-butyl group, 1,1-dimethylpropyl group, t-butoxycarbonyl group, tetrahydropyranyl group, tetrahydrofuranyl group, tetrahydrothiopyranyl group, tetrahydrothiofuranyl group, and the like are preferable.

The acid-dissociable group-containing resin (B) may contain one or more acid-dissociable groups.

[0106]

The proportion of the amount of the acid-dissociable group introduced into the acid-dissociable group-containing resin (B) (the proportion of the amount of the number of acid-dissociable groups in the total number of acidic functional groups and acid-dissociable groups in the acid-dissociable group-containing resin (B)) is preferably 5-100%, and still more preferably 10-100%, although the amount varies depending on the type of acid-dissociable group and the type of resin into which the acid-dissociable group is introduced.

Various types of structure for the acid-dissociable group-containing resin (B) may be used without any restrictions as long as the above properties can be obtained.

Preferable structures include a poly(p-hydroxystyrene) in which part or all of the hydrogen atoms in the phenolic hydroxyl groups are replaced by acid-dissociable groups, a copolymer of p-hydroxystyrene and/or p-hydroxy- α -methylstyrene and (meth)acrylic acid in which part or all of the hydrogen atoms in the phenolic hydroxyl groups and/or carboxyl groups are replaced by acid-dissociable groups, and the like.

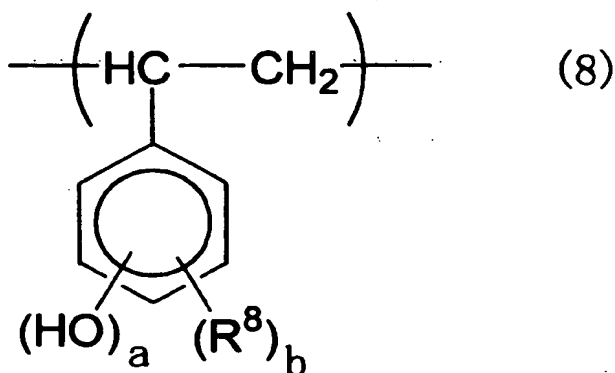
[0107]

Also, the structure of the acid-dissociable group-containing resin (B) can be appropriately determined according to the type of radiation employed.

As a preferable example of an acid-dissociable group-containing resin (B) especially suitable for a radiation-sensitive resin composition using a KrF excimer laser, an alkali insoluble or scarcely soluble resin having at least one recurring unit of the following formula (8) (hereinafter referred to as "recurring unit (8)") and at least one recurring unit (8), wherein the phenolic hydroxyl group is protected by an acid-dissociable group can be given (such an acid-dissociable group-containing resin is hereinafter referred to as "resin (B1)"). The resin (B1) may be suitably used in radiation-sensitive resin compositions for use with other radiations such as ArF excimer laser, F₂ excimer laser, and electron beams.

[0108]

[Chemical Formula 28]



wherein R⁸ represents a hydrogen atom or monovalent organic group, two or more R⁸s, if present, may be the same or different, and a and b each represent an integer of 1-3.

[0109]

As the recurring unit (8), units wherein the non-aromatic double bond of p-hydroxystyrene is cleaved are particularly preferable.

The resin (B1) may contain at least one other recurring unit.

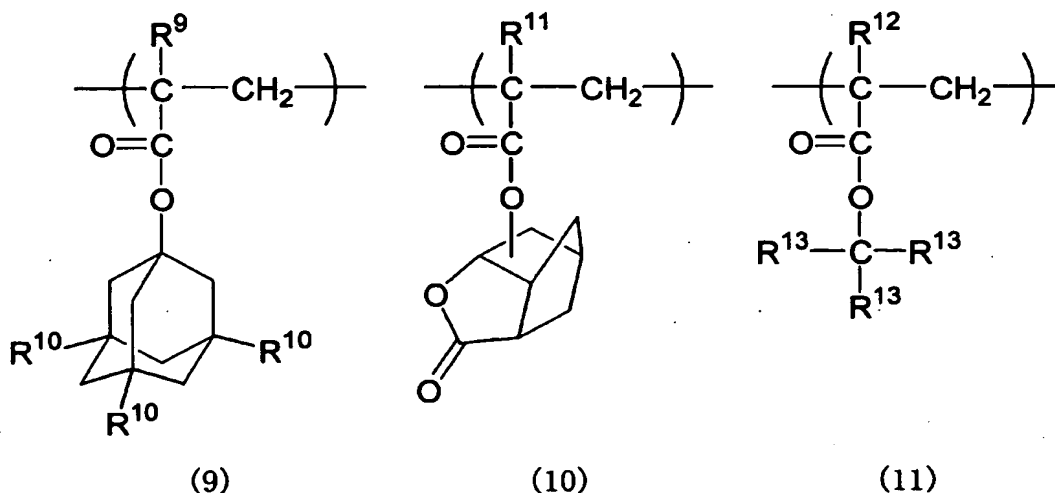
As examples of the other recurring unit, units obtained by cleavage of a polymerizable unsaturated bond of vinyl aromatic compounds such as styrene; (meth)acrylic esters such as t-butyl (meth)acrylate, adamantyl (meth)acrylate, and 2-methyladamantyl (meth)acrylate; and the like can be given.

[0110]

As a preferable example of an acid-dissociable group-containing resin (B) especially suitable for a radiation-sensitive resin composition using an ArF excimer laser, an alkali insoluble or scarcely soluble resin having at least one recurring unit selected from the group consisting of the recurring unit of the following formula (9) (hereinafter referred to as "recurring unit (9)") and the recurring unit of the following formula (10) (hereinafter referred to as "recurring unit (10)"), and at least one recurring unit of the following formula (11) (hereinafter referred to as "recurring unit (11)") can be given. This resin is hereinafter referred to as "resin (B2)". The resin (B2) may be suitably used in radiation-sensitive resin compositions for use with other radiations such as KrF excimer laser, F₂ excimer laser, and electron beams.

[0111]

[Chemical Formula 29]



[0112]

wherein R^9 , R^{11} , and R^{12} individually represent a hydrogen atom or a methyl group; R^{10} individually represents a hydrogen atom, hydroxyl group, cyano group, or $-COOR^{14}$,

5 wherein R^{14} is a hydrogen atom, a linear or branched alkyl group having 1-4 carbon atoms, or a cyclic alkyl group having 3-20 carbon atoms; R^{13} individually represents a monovalent alicyclic hydrocarbon group having 4-20 carbon atoms or a derivative thereof or a linear or branched alkyl group having 1-4 carbon atoms, provided that at least one R^{13} is the alicyclic hydrocarbon group or a derivative thereof or any two R^{13} groups form, in combination and together with the carbon atom to which these groups bond, a divalent alicyclic hydrocarbon group having 4-20 carbon atoms or a derivative thereof, and the remaining R^{13} group is a linear or branched alkyl group having 1-4 carbon atoms or a monovalent alicyclic hydrocarbon group having 4-20 carbon atoms or a derivative thereof.

15 [0113]

As preferable examples of the recurring unit (9), 3-hydroxyadamantan-1-yl (meth)acrylate, 3,5-dihydroxyadamantan-1-yl (meth)acrylate, 3-cyanoadamantan-1-yl (meth)acrylate, 3-carboxyladamantan-1-yl (meth)acrylate, 3,5-dicarboxyadamantan-1-yl (meth)acrylate, 3-carboxy-5-hydroxyadamantan-1-yl (meth)acrylate, 20 3-methoxycarbonyl-5-hydroxyadamantan-1-yl (meth)acrylate, and the like can be given.

The recurring units (9) and recurring units (10) may be present in the resin (B2) either individually or in combinations of two or more.

[0114]

As preferable examples of the recurring unit (11), 1-methyl-1-cyclopentyl (meth)acrylate, 1-ethyl-1-cyclopentyl (meth)acrylate, 1-methyl-1-cyclohexyl (meth)acrylate, 1-ethyl-1-cyclohexyl (meth)acrylate, 2-methyladamantan-2-yl (meth)acrylate, 2-ethyladamantan-2-yl (meth)acrylate, 2-n-propyladamantan-2-yl

(meth)acrylate, 2-i-propyladamantan-2-yl (meth)acrylate, 2-methyladamantan-2-yl (meth)acrylate, 1-(adamantan-1-yl)-1-methylethyl (meth)acrylate, and the like can be given.

[0115]

5 The resin (B2) may contain at least one other recurring unit.

As examples of the other recurring units, (meth)acrylic esters such as

7-oxo-6-oxabicyclo[3.2.1]octan-4-yl (meth)acrylate,

5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl (meth)acrylate, 2-oxotetrahydropyran-4-yl

(meth)acrylate, 4-methyl-2-oxotetrahydropyran-4-yl (meth)acrylate,

10 5-oxotetrahydrofuran-3-yl (meth)acrylate, 2-oxotetrahydrofuran-3-yl (meth)acrylate,

5-oxotetrahydrofuran-2-yl methyl (meth)acrylate, and

3,3-dimethyl-5-oxotetrahydrofuran-2-yl methyl (meth)acrylate; an unsaturated amide

compound such as (meth)acrylamide, N,N-dimethyl(meth)acrylamide, crotonamide,

maleinamide, fumaramide, mesaconamide, citraconamide, and itaconamide; unsaturated

15 carboxylic anhydrides such as maleic anhydride and itaconic anhydride;

mono-functional monomers such as bicyclo[2.2.1]hept-2-ene and the derivative thereof

and tetracyclo[6.2.1^{3,6}.0^{2,7}]dodec-3-ene and the derivative thereof; a polyfunctional

monomers such as methyleneglycol di(meth)acrylate, ethyleneglycol di(meth)acrylate,

2,5-dimethyl-2,5-hexanediol di(meth)acrylate, 1,2-adamantandiol di(meth)acrylate,

20 1,3-adamantandiol di(meth)acrylate, 1,4-adamantandiol di(meth)acrylate, and

tricyclodecanyl dimethylol di(meth)acrylate can be given.

[0116]

As a preferable example of the acid-dissociable group-containing resin (B)

especially suitable for a radiation-sensitive resin composition using an F₂ excimer laser,

25 an alkali insoluble or scarcely soluble polysiloxane having at least one recurring unit

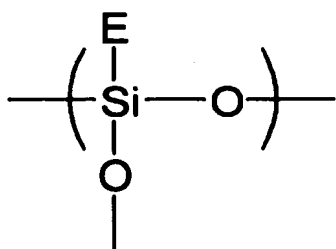
selected from the group consisting of the recurring unit of the following formula (12)

(hereinafter referred to as "recurring unit (12)") and the recurring unit of the following

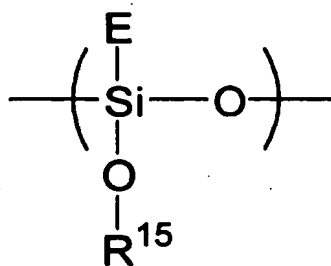
formula (13) (hereinafter referred to as “recurring unit (13)”) can be given (this resin is hereinafter referred to as “resin (B3)”). The resin (B3) may be suitably used in radiation-sensitive resin compositions for use with other radiations such as KrF excimer laser, ArF excimer laser, and electron beams.

5 [0117]

[Chemical Formula 30]



(12)



(13)

wherein E individually represents a monovalent organic group having an
10 acid-dissociable group and R^{15} represents a substituted or unsubstituted, linear, branched, or cyclic monovalent hydrocarbon group having 1-20 carbon atoms.

[0118]

E is preferably a group having a structure wherein an acid-dissociable group is bonded with a group having a cyclic structure.

15 As a preferable group having the cyclic structure, a group having an alicyclic ring structure derived from a cycloalkane having 3-8 carbon atoms, tricyclodecane, tetracyclodecane, adamantane, and the like, and a group having a halogenated aromatic ring structure having 6-20 carbon atoms can be given.

[0119]

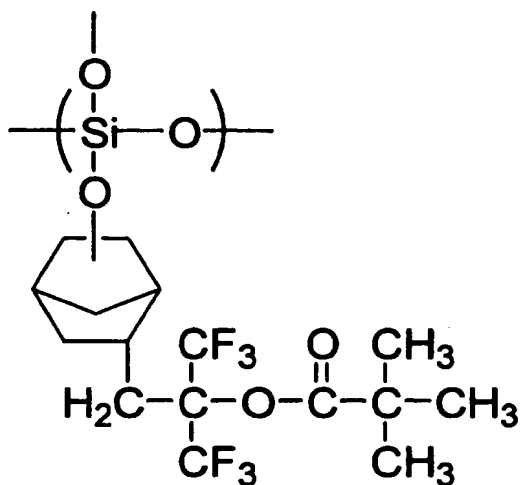
20 As the resin (B3), a resin having the recurring unit (12) is preferable.

As specific preferable examples of the recurring unit (12), the recurring units of

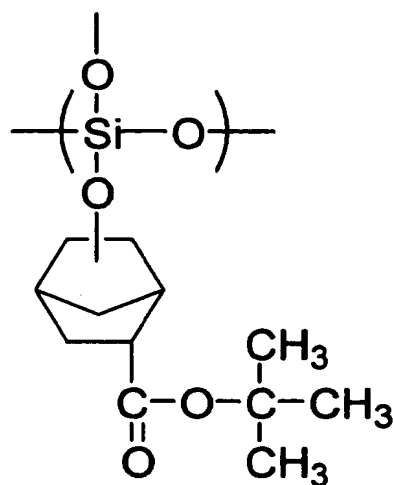
the following formulas (12-1)-(12-4) can be given.

[0120]

[Chemical Formula 31]



(12-1)

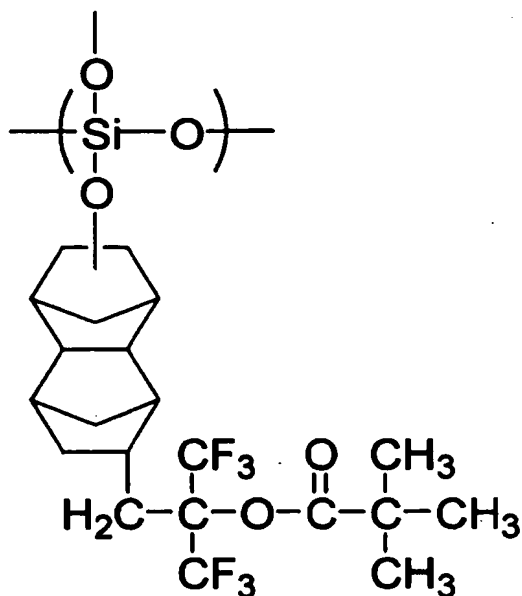


(12-2)

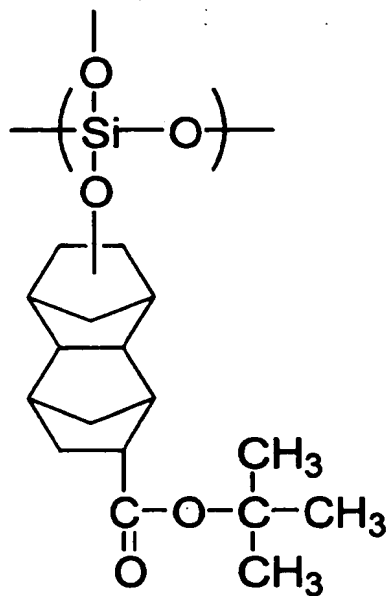
5

[0121]

[Chemical Formula 32]



(12-3)



(12-4)

[0122]

The recurring units (12) and recurring units (13) may be present in the resin (B3)

5 either individually or in combinations of two or more.

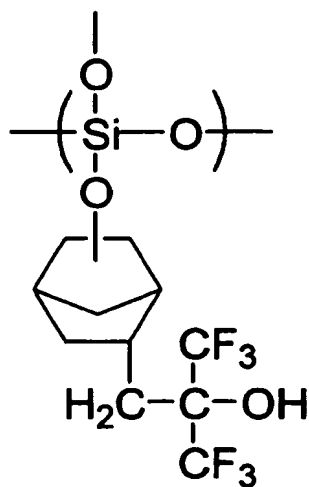
The resin (B3) may contain at least one other recurring unit.

As preferable examples of the other recurring unit, units obtained from hydrolysis of alkylalkoxysilanes such as methyltrimethoxysilane, methyltriethoxysilane, ethyltrimethoxysilane, and ethyltriethoxysilane, and the recurring units of the following

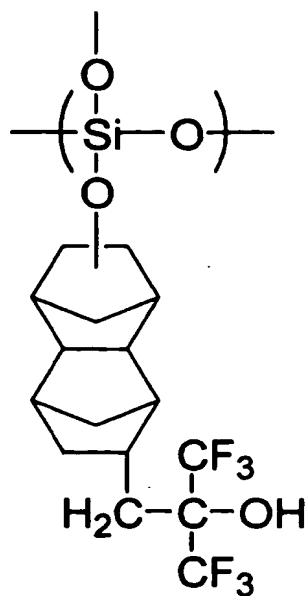
10 formulas (14-1)-(14-4) can be given.

[0123]

[Chemical Formula 33]



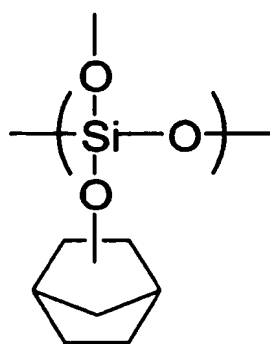
(14-1)



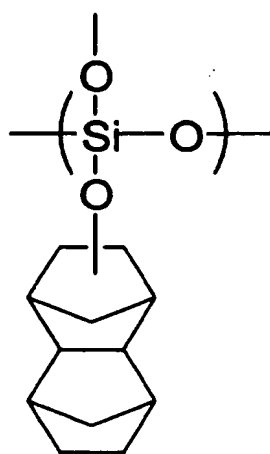
(14-2)

[0124]

[Chemical Formula 34]



(14-3)



(14-4)

[0125]

The resin (B3) can be prepared by polycondensation of a silane compound containing an acid-dissociable group or by introducing an acid-dissociable group into a previously prepared polysiloxane.

When polycondensing the acid-dissociable group-containing silane compound, an acidic catalyst is preferably used as the catalyst, and after polycondensation of the silane compound in the presence of the acidic catalyst, a further reaction is preferably continued in the presence of a basic catalyst.

[0126]

As examples of the acidic catalyst, inorganic acids such as hydrochloric acid, sulfuric acid, nitric acid, boric acid, phosphoric acid, titanium tetrachloride, zinc chloride, and aluminium chloride and organic acids such as formic acid, acetic acid, n-propionic acid, butyric acid, valeric acid, oxalic acid, malonic acid, succinic acid, maleic acid, fumaric acid, adipic acid, phthalic acid, terephthalic acid, acetic anhydride, maleic anhydride, citric acid, benzenesulfonic acid, p-toluenesulfonic acid, and methanesulfonic acid can be given.

Of these acidic catalysts, hydrochloric acid, sulfuric acid, acetic acid, oxalic acid, malonic acid, maleic acid, fumaric acid, acetic anhydride, maleic anhydride, and the like are preferable.

These acidic catalysts may be used either individually or in combination of two or more.

[0127]

As examples of the basic catalyst, inorganic bases such as lithium hydroxide, sodium hydroxide, potassium hydroxide, calcium hydroxide, barium hydroxide, sodium hydrogencarbonate, potassium hydrogencarbonate, sodium carbonate, and potassium carbonate and organic bases such as triethylamine, tri-n-propylamine, tri-n-butylamine, and pyridine can be given.

These basic catalysts may be used either individually or in combination of two or

more.

[0128]

When the acid-dissociable group-containing resin (B) is prepared by a reaction or reactions comprising the polymerization of a polymerizable unsaturated monomer, a
5 branched structure may be introduced into the acid-dissociable group-containing resin by a unit derived from a polyfunctional monomer having two or more polymerizable unsaturated bonds and/or by an acetal crosslinking group. Introduction of the branched structure improves the heat resistance of the acid-dissociable group-containing resin (B).

10 The amount of the branched structure introduced into the acid-dissociable group-containing resin (B) is preferably 10 mol% or less of the total amount of recurring units, although such an amount varies depending on the type of branched structure and the type of acid-dissociable group-containing resin into which the branched structure is introduced.

15 [0129]

The molecular weight of the acid-dissociable group-containing resin (B) may be appropriately selected without any restrictions. The polystyrene-reduced weight molecular weight (hereinafter referred to as "Mw") of the acid-dissociable group-containing resin (B) determined by gel permeation chromatography (GPC) is
20 usually 1,000-500,000, preferably 2,000-400,000, and still more preferably 3,000-300,000.

The Mw of the acid-dissociable group-containing resin (B) not having a branched structure is preferably 1,000-150,000, and particularly preferably 3,000-100,000. The Mw of the acid-dissociable group-containing resin (B) having a branched structure is
25 preferably 5,000-500,000, and particularly preferably 8,000-300,000. The resist obtained from the acid-dissociable group-containing resin (B) having an Mw in the above range possesses excellent development characteristics.

[0130]

The ratio of Mw to the polystyrene-reduced number molecular weight (hereinafter referred to as "Mn") determined by GPC (Mw/Mn) of the acid-dissociable group-containing resin (B) can be appropriately selected without any restrictions, and is usually 1-10, preferably 1-8, and particularly preferably 1-5. The resist obtained from the acid-dissociable group-containing resin (B) having a Mw/Mn in the above range possesses excellent resolution performance.

[0131]

There are no restrictions to the method for manufacturing the acid-dissociable group-containing resin (B). As examples of the method for manufacturing, a method of introducing one or more acid-dissociable groups into an acidic functional group of an alkali-soluble resin which has previously been manufactured, a method of polymerizing one or more polymerizable unsaturated monomers having an acid-dissociable group, optionally together with other polymerizable unsaturated monomers, a method of polycondensing one or more polycondensable components having an acid-dissociable group, optionally together with other polycondensable components, and the like can be given.

[0132]

The polymerization of the polymerizable unsaturated monomers and the polymerization of the one or more polymerizable unsaturated monomers possessing an acid-dissociable group in the manufacture of the alkali soluble resin is carried out by block polymerization, solution polymerization, precipitation polymerization, emulsion polymerization, suspension polymerization, block-suspension polymerization, or the like using an appropriate polymerization initiator or catalyst such as a radical polymerization initiator, anionic polymerization catalyst, coordinating anionic polymerization catalyst, cationic polymerization catalyst, or the like according to the type of polymerizable unsaturated monomer or reaction media.

The polycondensation of the one or more polycondensable components having an acid-dissociable group is preferably carried out in the presence of an acidic catalyst using an aqueous medium or a mixture of water and a hydrophilic solvent.

[0133]

5 The amount of the acid generator (A) used in the positive-tone radiation-sensitive resin composition of the present invention can be appropriately selected depending on the desired properties of the resist. The acid generator (A) is preferably used in an amount of 0.001-70 parts by weight, more preferably 0.01-50 parts by weight, and particularly preferably 0.1-20 parts by weight for 100 parts by weight of the
10 acid-dissociable group-containing resin (B). Using the acid generator (A) in an amount of 0.001 parts by weight or more prevents deterioration of the sensitivity and resolution of the resist. Also, using the acid generator (A) in an amount of 70 parts by weight or less prevents deterioration of the applicability and pattern shape of the resist.

[0134]

15 Acid diffusion controller

An acid diffusion controller is preferably added to the positive-tone radiation-sensitive resin composition of the present invention. The acid diffusion controller controls diffusion of an acid generated from the acid generator (A) upon exposure in the resist film and prevents unfavorable chemical reactions in the
20 unexposed region. Addition of the acid diffusion controller further improves storage stability of the resulting radiation-sensitive resin composition and resolution of the resist. Moreover, addition of the acid diffusion controller prevents the line width of the resist pattern from changing due to changes in the post-exposure delay (PED) between exposure and development, whereby a radiation-sensitive resin composition with
25 remarkably superior process stability can be obtained.

[0135]

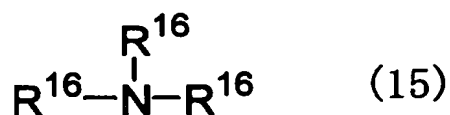
As the acid diffusion controller, nitrogen-containing organic compounds of which

the basicity does not change due to exposure or heat treatment during formation of a resist pattern are preferable.

As examples of the nitrogen-containing organic compounds, a compound shown by the following formula (15) (hereinafter referred to as “nitrogen-containing compound (I)”), a diamino compound having two nitrogen atoms in the molecule (hereinafter referred to as “nitrogen-containing compound (II)”), a polyamino compound or polymer having three or more nitrogen atoms in the molecule (hereinafter referred to as “nitrogen-containing compound (III)”), an amide group-containing compound, urea compound, nitrogen-containing heterocyclic compound, and the like can be given,

[0136]

[Chemical Formula 35]



wherein R¹⁶ individually represents a hydrogen atom, alkyl group, aryl group, or aralkyl group which may be substituted or unsubstituted.

[0137]

Given as examples of the substituted or unsubstituted alkyl group represented by R¹⁶ in the above formula (15) are groups having 1-15 carbon atoms and preferably 1-10 carbon atoms including specific examples such as a methyl group, ethyl group, n-propyl group, i-propyl group, n-butyl group, i-butyl group, sec-butyl group, t-butyl group, n-pentyl group, neopentyl group, n-hexyl group, n-heptyl group, n-octyl group, n-ethylhexyl group, n-nonyl group, and n-decyl group.

[0138]

Given as examples of the substituted or unsubstituted aryl group represented by R¹⁶ are groups having 6-12 carbon atoms including specific examples such as a phenyl

group, tolyl group, xylyl group, cumenyl group, and 1-naphthyl group.

Given as examples of the substituted or unsubstituted aralkyl group represented by R¹⁶ are groups having 7-19 carbon atoms and preferably 7-13 carbon atoms including specific examples such as a benzyl group, α -methylbenzyl group, phenethyl group, and 1-naphthylmethyl group.

[0139]

Examples of the nitrogen-containing compounds (I) include monoalkylamines such as n-hexylamine, n-heptylamine, n-octylamine, n-nonylamine, and n-decylamine; dialkylamines such as di-n-butylamine, di-n-pentylamine, di-n-hexylamine, di-n-heptylamine, di-n-octylamine, di-n-nonylamine, and di-n-decylamine; trialkylamines such as triethylamine, tri-n-propylamine, tri-n-butylamine, tri-n-pentylamine, tri-n-hexylamine, tri-n-heptylamine, tri-n-octylamine, tri-n-nonylamine, and tri-n-decylamine; aromatic amines such as aniline, N-methylaniline, N,N-dimethylaniline, 2-methylaniline, 3-methylaniline, 4-methylaniline, 4-nitroaniline, diphenylamine, triphenylamine, and 1-naphthylamine; and alkanolamines such as ethanolamine, diethanolamine, and triethanolamine.

[0140]

Examples of the nitrogen-containing compounds (II) include ethylenediamine, N,N,N',N'-tetramethylethylenediamine, tetramethylenediamine, hexamethylenediamine, N,N,N',N'-tetrakis(2-hydroxyethyl)ethylenediamine, N,N,N',N'-tetrakis(2-hydroxypropyl)ethylenediamine, 4,4'-diaminodiphenylmethane, 4,4'-diamino diphenyl ether, 4,4'-diaminobenzophenone, 4,4'-diaminodiphenylamine, 2,2'-bis(4-aminophenyl)propane, 2-(3-aminophenyl)-2-(4-aminophenyl)propane, 2-(4-aminophenyl)-2-(3-hydroxyphenyl)propane, 2-(4-aminophenyl)-2-(4-hydroxyphenyl)propane, 1,4-bis[1-(4-aminophenyl)-1-methylethyl]benzene, and 1,3-bis[1-(4-aminophenyl)-1-methylethyl]benzene.

As examples of the nitrogen-containing compound (III), polyethyleneimine, polyallylamine, a polymer of dimethylaminoethylacrylamide, and the like can be given.

[0141]

Examples of compounds containing an amide group include formamide, N-methylformamide, N,N-dimethylformamide, acetamide, N-methylacetamide, N,N-dimethylacetamide, propionamide, benzamide, pyrrolidone, N-methylpyrrolidone, and the like.

Examples of urea compounds include urea, methylurea, 1,1-dimethylurea, 1,3-dimethylurea, 1,1,3,3-tetramethylurea, 1,3-diphenylurea, and tributylthiourea.

[0142]

Examples of the nitrogen-containing heterocyclic compounds include imidazoles such as imidazole, benzimidazole, 2-methylimidazole, 4-methylimidazole, 1,2-dimethylimidazole, 2-phenylimidazole, 4-phenylimidazole, 4-methyl-2-phenylimidazole, and 2-phenylbenzimidazole; pyridines such as pyridine, 2-methylpyridine, 4-methylpyridine, 2-ethylpyridine, 4-ethylpyridine, 2-phenylpyridine, 4-phenylpyridine, N-methyl-4-phenylpyridine, nicotine, nicotinic acid, nicotinamide, quinoline, 8-oxyquinoline, and acridine; pyrazine, pyrazole, pyridazine, quinoxaline, purine, pyrrolidine, piperidine, 1-piperidineethanol, 2-piperidineethanol, 3-piperidine-1,2-propanediol, morpholine, 4-methylmorpholine, piperazine, 1,4-dimethylpiperazine, and 1,4-diazabicyclo[2.2.2]octane.

[0143]

A compound having an acid-dissociable group can also be used as the nitrogen-containing organic compound.

As examples of the nitrogen-containing organic compound having an acid-dissociable group, N-(t-butoxycarbonyl)piperidine, N-(t-butoxycarbonyl)imidazole, N-(t-butoxycarbonyl)benzimidazole, N-(t-butoxycarbonyl)-2-phenylbenzimidazole, N-(t-butoxycarbonyl)dioctylamine, N-(t-butoxycarbonyl)diethanolamine,

N-(t-butoxycarbonyl)dicyclohexylamine, and N-(t-butoxycarbonyl)diphenylamine can be given.

[0144]

Of these nitrogen-containing organic compounds, the nitrogen-containing
5 compounds (I), nitrogen-containing compounds (II), nitrogen-containing heterocyclic compounds, and the like are preferable.

The acid diffusion controller may be used either individually or in combination of two or more.

[0145]

10 The amount of the acid diffusion controller to be added is preferably 15 parts by weight or less, more preferably 0.001-10 parts by weight, and particularly preferably 0.005-5 parts by weight for 100 parts by weight of the acid-dissociable group-containing resin (B). Incorporating the acid diffusion controller in an amount of 0.001 parts by weight or more prevents deterioration of the pattern shape and size
15 fidelity as a resist. Also, incorporating the acid diffusion controller in an amount of 15 parts by weight or less improves the sensitivity as a resist and improves the developability of the exposure area.

[0146]

Dissolution controller

20 A dissolution controller that improves the solubility in an alkaline developer by the action of an acid may be added to the positive-tone radiation-sensitive resin composition of the present invention.

As examples of such a dissolution controller, compounds having an acid functional group such as a phenolic hydroxyl group, carboxyl group, and sulfonic group,
25 compounds in which the hydrogen atom in the acidic functional group is replaced by an acid-dissociable group, and the like can be given.

These dissolution controllers may be used either individually or in combination of

two or more. The proportion of the dissolution controllers to be added is 10 parts by weight or less, and preferably 5 parts by weight or less for 100 parts by weight of the total resin component in the radiation-sensitive resin composition.

[0147]

5 Surfactant

A surfactant that improves applicability, striation, developability, and the like may be added to the positive-tone radiation-sensitive resin composition of the present invention.

As the surfactants, any of anionic surfactants, cationic surfactants, nonionic
10 surfactants, and ampholytic surfactants may be used. Of these, nonionic surfactants are preferable.

As examples of nonionic-type surfactants, polyoxyethylene higher alkyl ethers, polyoxyethylene higher alkyl phenyl ethers, higher fatty acid diesters of polyethylene glycol, commercially available products such as KP (manufactured by Shin-Etsu
15 Chemical Co., Ltd.), Polyflow (manufactured by Kyoeisha Chemical Co., Ltd.), EFTOP (manufactured by Tohkem Products Corporation), MEGAFAC (manufactured by Dainippon Ink and Chemicals, Inc.), Fluorad (manufactured by Sumitomo 3M, Ltd.), Asahi Guard, Surflon (manufactured by Asahi Glass Co., Ltd.), and the like can be given.

20 These surfactants may be used either individually or in combination of two or more. The proportion of the surfactants to be added is 2 parts by weight or less, and preferably 1.5 parts by weight or less, as an effective component, for 100 parts by weight of the total resin components in the radiation-sensitive resin composition.

[0148]

25 Photosensitizer

Sensitizers can be added to the positive-tone radiation-sensitive resin composition of the present invention. These sensitizers absorb radiation energy and transmit the

energy to the acid generator (A), thereby increasing the amount of an acid to be generated upon exposure and improving the apparent sensitivity of the radiation-sensitive resin composition.

As examples of preferable sensitizers, acetophenones, benzophenones, naphthalenes, biacetyl, eosine, rose bengale, pyrenes, anthracenes, phenothiazines, and the like can be given.

These sensitizers may be used either individually or in combinations of two or more. The proportion of the sensitizers to be added is 50 parts by weight or less, and preferably 30 parts by weight or less for 100 parts by weight of the total resin component in the radiation-sensitive resin composition.

[0149]

Other additives

Other additives may be added to the positive-tone radiation-sensitive resin composition of the present invention, as required, to the extent that does not impair the effects of the present invention. Examples of such additives include dyes, pigments, adhesion adjuvants, halation inhibitors, preservatives, defoaming agents, and shape improvers. Specific additives include 4-hydroxy-4'-methylchalcone, and the like.

Addition of a dye or a pigment visualizes a latent image in the exposed area, thereby decreasing the effects of halation during exposure. Use of an adhesion improver improves adhesion to the substrates.

[0150]

Preparation of composition solution

The positive-tone radiation-sensitive resin composition of the present invention is usually prepared as a composition solution by dissolving the components in a solvent to obtain a homogeneous solution and, optionally, filtering the solution through a filter with a pore size of about 0.2 μm .

[0151]

Ethers, esters, ether esters, ketones, ketone esters, amides, amide esters, lactams, lactones, and (halogenated) hydrocarbons are given as examples of the solvent which can be used here. Specific examples are ethylene glycol monoalkyl ethers, diethylene glycol dialkyl ethers, propylene glycol monoalkyl ethers, propylene glycol dialkyl
 5 ethers, ethylene glycol monoalkyl ether acetates, propylene glycol monoalkyl ether acetates, acetates, hydroxyacetates, lactates, alkoxyacetates, (non)cyclic ketones, acetoacetates, pyruvates, propionates, N,N-dialkylformamides, N,N-dialkylacetamides, N-alkylpyrrolidones, γ -lactones, (halogenated) aliphatic hydrocarbons, and (halogenated) aromatic hydrocarbons.

10 [0152]

More specifically, such solvents include ethylene glycol monomethyl ether, ethylene glycol monoethyl ether, ethylene glycol mono-n-propyl ether, ethylene glycol mono-n-butyl ether, diethylene glycol dimethyl ether, diethylene glycol diethyl ether, diethylene glycol di-n-propyl ether, diethylene glycol di-n-butyl ether, ethylene glycol
 15 monomethyl ether acetate, ethylene glycol monoethyl ether acetate, propylene glycol monomethyl ether acetate, propylene glycol monoethyl ether acetate, propylene glycol mono-n-propyl ether acetate, isopropenyl acetate, isopropenyl propionate, toluene, xylene, methyl ethyl ketone, cyclohexanone, 2-heptanone, 3-heptanone, 4-heptanone, ethyl 2-hydroxypropionate, ethyl 2-hydroxy-2-methylpropionate, ethyl ethoxyacetate,
 20 ethyl hydroxyacetate, methyl 2-hydroxy-3-methylbutyrate, methyl lactate, ethyl lactate, n-propyl lactate, i-propyl lactate, 3-methoxybutyl acetate, 3-methyl-3-methoxybutyl acetate, 3-methyl-3-methoxybutyl propionate, 3-methyl-3-methoxybutyl butyrate, ethyl acetate, n-propyl acetate, n-butyl acetate, methyl acetoacetate, ethyl acetoacetate, methyl 3-methoxypropionate, ethyl 3-methoxypropionate, methyl 3-ethoxypropionate,
 25 ethyl 3-ethoxypropionate, N-methylpyrrolidone, N,N-dimethylformamide, and N,N-dimethylacetamide.

[0153]

Of these solvents, propylene glycol monoalkyl ether acetates, 2-heptanone, lactates, 2-hydroxypropionates, 3-alkoxypropionates, and the like are desirable to ensure excellent uniformity of the film surface during application.

These solvents may be used either individually or in combination of two or more.

5 [0154]

One or more solvents with a high boiling point may optionally be added to the solvent. Examples of such solvents with a high boiling point include benzyl ethyl ether, di-n-hexyl ether, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, acetonylacetone, isophorone, caproic acid, caprylic acid, 1-octanol, 10 1-nonanol, benzyl alcohol, benzyl acetate, ethyl benzoate, diethyl oxalate, diethyl maleate, γ -butyrolactone, ethylene carbonate, propylene carbonate, and ethylene glycol monophenyl ether acetate.

These other solvents may be used either individually or in combination of two or more.

15 The proportion of the other solvents to be added is 50 wt% or less, and preferably 30 wt% or less of the total amount of solvents used.

[0155]

The solvents are used in a total amount to make the total solid content of the solution composition usually 5-50 wt%, preferably 10-50 wt%, more preferably 10-40 20 wt%, and particular preferably 10-30 wt%. An optimal amount is 10-25 wt%. The total solid content in the above ranges is desirable to ensure excellent uniformity of the film surface during application.

[0156]

Formation of resist pattern

25 A resist pattern is formed from the positive-tone radiation-sensitive resin composition of the present invention by applying the composition solution thus prepared to, for example, substrates such as a silicon wafer or a wafer coated with

aluminum using an appropriate application method such as rotational coating, cast coating, and roll coating to form a resist film. Then, after optional heat treatment (hereinafter referred to as "PB"), the resist film is exposed to radiation through a mask with a prescribed pattern.

5 As radiation that can be used here, far ultraviolet rays such as a bright line spectrum of a mercury lamp (wavelength: 254 nm), KrF excimer laser (wavelength: 248 nm), ArF excimer laser (wavelength: 193 nm), F₂ excimer laser (wavelength: 157 nm), and EUV (wavelength: 13 nm); X-rays such as synchrotron radiation, charged particle rays such as electron beams, and the like can be used according to the types of acid
10 generator (A).

Of these, far ultraviolet rays and charged particle rays are preferable. Particular preferable radiations are a KrF excimer laser (wavelength: 248 nm), ArF excimer laser (wavelength: 193 nm), and electron beams.

The exposure conditions such as the dose of radiation are appropriately
15 determined according to the composition of the positive-tone radiation-sensitive resin composition, types of additives, and the like.

When forming a resist pattern, post exposure bake (hereinafter called "PEB") which is a heat treatment after exposure is preferable to increase apparent sensitivity of the resist.

20 PEB is performed at a temperature of 30-200°C, and preferably 50-150°C, although the temperature varies depending on the composition of the radiation-sensitive resin composition, types of additives, and the like.

[0157]

The resist film after exposure is developed in an alkaline developer to form a
25 predetermined resist pattern.

As the alkaline developer, an alkaline aqueous solution in which one or more alkaline compounds such as an alkaline metal hydroxide, aqueous ammonia,

alkylamines, alkanolamines, heterocyclic amines, tetraalkylammonium hydroxides, choline, 1,8-diazabicyclo[5.4.0]-7-undecene, and 1,5-diazabicyclo[4.3.0]-5-nonene are dissolved is used. An aqueous solution of tetraalkylammonium hydroxide is a particularly preferable alkaline developer.

- 5 The concentration of the alkaline aqueous solution is preferably 10 wt% or less, more preferably 1-10 wt%, and particularly preferably 2-5 wt%. The concentration of the alkaline aqueous solution less than 10 wt% prevents dissolution of an unexposed area in the developer.

- The addition of an appropriate amount of a surfactant to the alkaline aqueous
10 solution is desirable to increase wettability of the resist to the developer.

 After development using the alkaline aqueous solution developer, the resist film is generally washed with water and dried.

[0158]

[Examples]

- 15 The embodiments of the present invention are described in more detail by examples. However, these examples should not be construed as limiting the present invention.

<Synthesis of acid generator>

Synthesis Example 1

- 20 26.7 g of 4-fluorophenyl.diphenylsulfonium trifluoromethanesulfonate was dissolved in 200 g of dichloromethane in a reaction flask, in which the atmosphere was replaced with nitrogen. 200 g of 10 wt% sodium hydroxide aqueous solution, 5.8 g of tetra-n-butylammonium bromide, and 6.71 g of n-butanethiol were added. The mixture was stirred for 30 minutes at room temperature and transferred to a separatory
25 funnel, shaken, and allowed to stand still. The water layer was removed. After the addition of 300 ml of distilled water, the mixture was shaken again and allowed to stand still. The water layer was removed. The resulting dichloromethane solution was

dried with anhydrous magnesium sulfate, filtered, and dried using an evaporator. After evaporating dichloromethane from the dried solution, the residual liquid was dried under reduced pressure to obtain 17.8 g of 4-n-butylthiophenyl.diphenylsulfonium trifluoromethanesulfonate.

5 [0159]

17.8 g of the resulting 4-n-butylthiophenyl.diphenylsulfonium trifluoromethanesulfonate was dissolved in 100 g of methanol in a reaction flask. The mixture was fed into an ion-exchange chromatography (an ion-exchange resin: Shephadex R-QAE A-25, packing amount: 60 g, manufactured by Aldrich Corporation) to replace trifluoromethanesulfonate anions with chlorine ions. After evaporating methanol using an evaporator, the resulting residue was dissolved in 100 g of dichloromethane. 15 g of 30 wt% nonafluoro-n-butanesulfonic acid ammonium aqueous solution was added. The mixture was stirred for one hour at room temperature and transferred to a separatory funnel, shaken, and allowed to stand still. The water layer was removed. 100 ml of distilled water was added and the mixture was shaken again and allowed to stand still. The water layer was removed. The resulting dichloromethane solution was dried with anhydrous magnesium sulfate, filtered, and dried using an evaporator. After evaporating dichloromethane from the dried solution, the residual liquid was dried under reduced pressure to obtain 21.1 g of 4-n-butylthiophenyl.diphenylsulfonium nonafluoro-n-butanesulfonate.

[0160]

21.1 g of the resulting 4-n-butylthiophenyl.diphenylsulfonium nonafluoro-n-butanesulfonate was dissolved in 300 ml of methanol in a reaction flask. 18 g of 30 wt% hydrogen peroxide solution and 10.5 g of sodium tungstate dihydride were added. The mixture was stirred for 30 minutes at room temperature. The resulting methanol solution was condensed under reduced pressure and the residue was dissolved in 300 g of dichloromethane. The mixture was transferred to a separatory

funnel, shaken, and allowed to stand still. The water layer was removed. 100 ml of distilled water was added and the mixture was shaken again and allowed to stand still. The water layer was removed. The resulting dichloromethane solution was dried with anhydrous magnesium sulfate, filtered, and dried using an evaporator. After
5 evaporating dichloromethane from the dried solution, the residual liquid was dried under reduced pressure to obtain 14.4 g of 4-n-butylsulfonylphenyl.diphenylsulfonium nonafluoro-n-butanesulfonate. This compound is indicated as "acid generator (A-1)".

[0161]

Synthesis Example 2

10 26.7 g of 4-fluorophenyl.diphenylsulfonium trifluoromethanesulfonate was dissolved in 200 g of dichloromethane in a reaction flask, in which the atmosphere was replaced with nitrogen. 200 g of 10 wt% sodium hydroxide aqueous solution, 5.8 g of tetra-n-butylammonium bromide, and 6.71 g of cyclohexanethiol were added. The mixture was stirred for 30 minutes at room temperature and transferred to a separatory
15 funnel, shaken, and allowed to stand still. The water layer was removed. After the addition of 300 ml of distilled water, the mixture was shaken again and allowed to stand still. The water layer was removed. The resulting dichloromethane solution was dried with anhydrous magnesium sulfate, filtered, and dried using an evaporator. After
20 evaporating dichloromethane from the dried solution, the residual liquid was dried under reduced pressure to obtain 19.2 g of 4-cyclohexylthiophenyl.diphenylsulfonium trifluoromethanesulfonate.

[0162]

19.2 g of the resulting 4-cyclohexylthiophenyl.diphenylsulfonium trifluoromethane sulfonate was dissolved in 100 g of methanol in a reaction flask. The
25 mixture was put through an ion-exchange chromatography (an ion-exchange resin: Shephadex R-QAE A-25, 60g of filling amount, manufactured by Aldrich Chemical Co., Inc.) to replace trifluoromethanesulfonate anions with chlorine ions. After evaporating

methanol using an evaporator, the resulting residue was dissolved in 100 g of dichloromethane. 15 g of 30 wt% nonafluoro-n-butane sulfonic acid ammonium aqueous solution was added. The mixture was stirred for one hour at room temperature and transferred to a separatory funnel, shaken, and allowed to stand still.

- 5 The water layer was removed. 100 ml of distilled water was added and the mixture was shaken again and allowed to stand still. The water layer was removed. The resulting dichloromethane solution was dried with anhydrous magnesium sulfate, filtered, and dried using an evaporator. After evaporating dichloromethane from the dried solution, the residual liquid was dried under reduced pressure to obtain 20.5 g of
- 10 4-cyclohexylthiophenyl.diphenylsulfonium nonafluoro-n-butanesulfonate.

[0163]

- 20.5 g of the resulting 4-cyclohexylthiophenyl.diphenylsulfonium nonafluoromethane-n-butanesulfonate was dissolved in 300 ml of methanol in a reaction flask. 18 g of 30 wt% hydrogen peroxide solution and 10.5 g of sodium
- 15 tangstate dihydride were added. The mixture was stirred for 30 minutes at room temperature. The resulting methanol solution was condensed under reduced pressure and the residue was dissolved in 300 g of dichloromethane. The mixture was transferred to a separatory funnel, shaken, and allowed to stand still. The water layer was removed. 100 ml of distilled water was added and the mixture was shaken again
- 20 and allowed to stand still. The water layer was removed. The resulting dichloromethane solution was dried with anhydrous magnesium sulfate, filtered, and dried using an evaporator. After evaporating dichloromethane from the dried solution, the residual liquid was dried under reduced pressure to obtain 15.5 g of
- 4-cyclohexylsulfonylphenyl.diphenylsulfonium nonafluoro-n-butanesulfonate. This
- 25 compound is indicated as "acid generator (A-2)".

[0164]

Synthesis Example 3

26.7 g of tris(4-fluorophenyl)sulfonium trifluoromethanesulfonate was dissolved in 200 g of dichloromethane in a reaction flask, in which the atmosphere was replaced with nitrogen. 600 g of 10 wt% sodium hydroxide aqueous solution, 5.8 g of tetra-n-butylammonium bromide, and 20.1 g of n-butanethiol were added. The mixture was stirred for 30 minutes at room temperature and transferred to a separatory funnel, shaken, and allowed to stand still. The water layer was removed. After the addition of 300 ml of distilled water, the mixture was shaken again and allowed to stand still. The water layer was removed. The resulting dichloromethane solution was dried with anhydrous magnesium sulfate, filtered, and dried using an evaporator. After evaporating dichloromethane from the dried solution, the residual liquid was dried under reduced pressure to obtain 23.4 g of tris(4-n-butylthiophenyl)sulfonium trifluoromethanesulfonate.

[0165]

23.4 g of the resulting tris(4-n-butylthiophenyl)sulfonium trifluoromethanesulfonate was dissolved in 100 g of methanol in a reaction flask. The mixture was fed into an ion-exchange chromatography (an ion-exchange resin: Shephadex R-QAE A-25, packing amount: 180 g, manufactured by Aldrich Corporation) to replace trifluoromethanesulfonate anions with chlorine ions. After evaporating methanol using an evaporator, the resulting residue was dissolved in 300 g of dichloromethane. 45 g of 30 wt% ammonium nonafluoro-n-butanesulfonate aqueous solution was added. The mixture was stirred for one hour at room temperature and transferred to a separatory funnel, shaken, and allowed to stand still. The water layer was removed. 100 ml of distilled water was added and the mixture was shaken again and allowed to stand still. The water layer was removed. The resulting dichloromethane solution was dried with anhydrous magnesium sulfate, filtered, and dried using an evaporator. After evaporating dichloromethane from the dried solution, the residual liquid was dried under reduced pressure to obtain 25.2 g of

tris(4-n-butylthiophenyl)sulfonium nonafluoro-n-butanesulfonate.

[0166]

25.2 g of the resulting tris(4-n-butylthiophenyl)sulfonium
nonafluoro-n-butanesulfonate was dissolved in 300 ml of methanol in a reaction flask.
5 54 g of 30 wt% hydrogen peroxide solution and 30 g of sodium tungstate dihydride
were added. The mixture was stirred for 30 minutes at room temperature. The
resulting methanol solution was condensed under reduced pressure and the residue was
dissolved in 300 g of dichloromethane. The mixture was transferred to a separatory
funnel, shaken, and allowed to stand still. The water layer was removed. 100 ml of
10 distilled water was added and the mixture was shaken again and allowed to stand still.
The water layer was removed. The resulting dichloromethane solution was dried with
anhydrous magnesium sulfate, filtered, and dried using an evaporator. After
evaporating dichloromethane from the dried solution, the residual liquid was dried
under reduced pressure to obtain 20.1 g of tris(4-n-butylsulfonylphenyl)sulfonium
15 nonafluoro-n-butanesulfonate. This compound is indicated as "acid generator (A-3)".

[0167]

Mass spectrometry analysis

The acid generators (A-1) to (A-3) were subjected to mass spectrometry analysis
using JMS-AX505W mass spectrometer manufactured by JEOL Ltd. The analysis
20 conditions were as follows. The spectra obtained for cationic moieties of the acid
generators are shown in Figures 1-3.

Emitter current	: 5 mA (used gas: Xe)
Acceleration voltage	: 3.0 kV
10 N MULTI	: 1.3
25 Ionization method	: Fast atom bombardment (FAB) method
Detected ion	: cation (+)
Measured mass range	: 20-1,500 m/z

Scanning : 30 sec
 Resolving power : 1,500
 Matrix : 3-Nitrobenzyl alcohol
 [0168]

5 ¹H-NMR Analysis

The acid generators (A-1) to (A-3) were subjected to ¹H-NMR analysis using JNM-EX270 manufactured by JEOL Ltd. The solvent used for the measurement was deuterated chloroform. The spectra obtained are shown in Figures 4-6.

[0169]

10 <Synthesis of acid-dissociable group-containing resin (B)>

Acid-dissociable group-containing resins (B) were synthesized according to the following method.

Mw and Mn of the acid-dissociable group-containing resins (B) were measured by gel permeation chromatography (GPC) using GPC columns (manufactured by Tosoh Corp., G2000HXL x 2, G3000HXL x 1, G4000HXL x 1) under the following conditions.
 15 Flow rate: 1.0 ml/minute, eluate: tetrahydrofuran, column temperature: 40°C, standard reference material: monodispersed polystyrene

[0170]

Synthesis Example 4

20 101 g of p-acetoxystyrene, 5 g of styrene, 42 g of p-t-butoxystyrene, 6 g of azobisisobutyronitrile, and 1 g of t-dodecylmercaptan were dissolved in 160 g of propylene glycol monomethyl ether. The mixture was polymerized for 16 hours while maintaining the reaction temperature at 70°C in a nitrogen atmosphere. After polymerization, the reaction solution was added dropwise to a large quantity of
 25 n-hexane to coagulate and purify the resulting resin. After the addition of 150 g of propylene glycol monomethyl ether to the purified resin, 300 g of methanol, 80 g of triethylamine, and 15 g of water were added. The mixture was hydrolyzed for eight

hours while refluxing at the boiling point. After the reaction, the solvent and triethylamine were evaporated under reduced pressure. The resulting resin was dissolved in acetone and added dropwise to a large quantity of water to coagulate the resin. The resulting white powder was filtered and dried overnight at 50°C under reduced pressure.

The resin was found to have Mw and Mw/Mn of 16,000 and 1.7 respectively. The result of ¹³C-NMR analysis confirmed that the copolymerization molar ratio of p-hydroxystyrene, styrene, and p-t-butoxystyrene of the copolymer was 72:5:23. This resin is indicated as a "resin (B-1)".

[0171]

Synthesis Example 5

100 g of p-acetoxystyrene, 25 g of t-butyl acrylate, 18 g of styrene, 6 g of azobisisobutyronitrile, and 1 g of t-dodecylmercaptan were dissolved in 230 g of propylene glycol monomethyl ether. The mixture was polymerized for 16 hours while maintaining the reaction temperature at 70°C in a nitrogen atmosphere. After polymerization, the reaction solution was added dropwise to a large quantity of n-hexane to coagulate and purify the resulting resin. After the addition of 150 g of propylene glycol monomethyl ether to the purified resin, 300 g of methanol, 80 g of triethylamine, and 15 g of water were added. The mixture was hydrolyzed for eight hours while refluxing at the boiling point. After the reaction, the solvent and triethylamine were evaporated under reduced pressure. The resulting resin was dissolved in acetone and added dropwise to a large quantity of water to coagulate the resin. The resulting white powder was filtered and dried overnight at 50°C under reduced pressure.

The resin was found to have Mw and Mw/Mn of 11,500 and 1.6 respectively. The result of ¹³C-NMR analysis confirmed that the copolymerization molar ratio of p-hydroxystyrene, t-butyl acrylate, and styrene of the copolymer was 61:19:20. This

resin is indicated as a "resin (B-2)".

[0172]

Synthesis Example 6

97 g of p-acetoxystyrene, 51 g of p-t-butoxystyrene, 6 g of azobisisobutyronitrile,
 5 and 1 g of t-dodecylmercaptan were dissolved in 160 g of propylene glycol monomethyl
 ether. The mixture was polymerized for 16 hours while maintaining the reaction
 temperature at 70°C in a nitrogen atmosphere. After polymerization, the reaction
 solution was added dropwise to a large quantity of n-hexane to coagulate and purify the
 resulting resin. After the addition of 150 g of propylene glycol monomethyl ether to
 10 the purified resin, 300 g of methanol, 80 g of triethylamine, and 15 g of water were
 added. The mixture was hydrolyzed for eight hours while refluxing at the boiling
 point. After the reaction, the solvent and triethylamine were evaporated under reduced
 pressure. The resulting resin was dissolved in acetone and added dropwise to a large
 quantity of water to coagulate the resin. The resulting white powder was filtered and
 15 dried overnight at 50°C under reduced pressure.

The resin was found to have Mw and Mw/Mn of 16,500 and 1.7 respectively.
 The result of ¹³C-NMR analysis confirmed that the copolymerization molar ratio of
 p-hydroxystyrene and p-t-butoxystyrene of the copolymer was 67:33. This resin is
 indicated as a "resin (B-3)".

20 [0173]

Synthesis Example 7

25 g of poly(p-hydroxystyrene) was dissolved in 80 g of n-butyl acetate, and
 nitrogen gas was bubbled through the solution for 30 minutes. After the addition of 49
 g of di-t-butyl dicarbonate and 25 g of triethylamine as a catalyst, the mixture was
 25 reacted for seven hours at 60°C. After the reaction, n-butyl acetate was evaporated
 under reduced pressure. The resulting resin was dissolved in acetone and added
 dropwise to a large quantity of water to coagulate the resin. The resulting white

powder was filtered and dried overnight at 50°C under reduced pressure.

Mw and Mw/Mn of this resin were respectively 12,000 and 1.7. As a result of ¹³C-NMR analysis, the resin was found to have a structure in which 26 mol% of hydrogen atoms of a phenolic hydroxyl group in poly(p-hydroxystyrene) was replaced
5 by t-butoxycarbonyl groups. This resin is indicated as a “resin (B-4)”.

[0174]

Synthesis Example 8

25 g of poly(p-hydroxystyrene) was dissolved in 100 g of propylene glycol monomethyl acetate, and nitrogen gas was bubbled through the solution for 30 minutes.
10 After the addition of 4.8 g of ethyl vinyl ether and, as a catalyst, 1 g of pyridinium p-toluenesulfonate, the mixture was reacted for 12 hours at room temperature. The reaction solution was dropped into 1 wt% ammonium aqueous solution to coagulate the resin. The precipitated white powder was filtered and dried overnight at 50°C under reduced pressure.

15 Mw and Mw/Mn of this resin were respectively 13,000 and 1.7. As a result of ¹H-NMR analysis, the resin was found to have a structure in which 34 mol% of hydrogen atoms of a phenolic hydroxyl group in poly(p-hydroxystyrene) was replaced by ethoxyethyl groups. This resin is indicated as a “resin (B-5)”.

[0175]

20 Synthesis Example 9

25 g of a copolymer of 92:8 p-hydroxystyrene and p-t-butoxycarbonyloxystyrene was dissolved in 100 g of propylene glycol monomethyl acetate, and nitrogen gas was bubbled through the solution for 30 minutes. After the addition of 3.3 g of ethyl vinyl ether and, as a catalyst, 1 g of pyridinium p-toluenesulfonate, the mixture was reacted
25 for 12 hours at room temperature. The reaction solution was dropped into 1 wt% ammonium aqueous solution to coagulate the resin. The precipitated white powder was filtered and dried overnight at 50°C under reduced pressure.

Mw and Mw/Mn of this resin were respectively 13,000 and 1.8. As a result of ^{13}C -NMR analysis, the resin was found to have a structure in which 23 mol% of hydrogen atoms of the phenolic hydroxyl group in poly(p-hydroxystyrene) was replaced by 1-ethoxyethyl groups, and 8 mol% by t-butyl groups. This resin is indicated as a
5 “resin (B-6)”.

[0176]

Synthesis Example 10

25 g of a copolymer of 90:10 p-hydroxystyrene and p-t-butoxystyrene was dissolved in 100 g of propylene glycol monomethyl acetate, and nitrogen gas was
10 bubbled through the solution for 30 minutes. After the addition of 3.3 g of ethyl vinyl ether and, as a catalyst, 1 g of pyridinium p-toluenesulfonate, the mixture was reacted for 12 hours at room temperature. The reaction solution was dropped into 1 wt% ammonium aqueous solution to coagulate the resin. The precipitated white powder was filtered and dried overnight at 50°C under reduced pressure.

15 Mw and Mw/Mn of this resin were respectively 13,000 and 1.01. As a result of ^{13}C -NMR analysis, the resin was found to have a structure in which 23 mol% of hydrogen atoms of the phenolic hydroxyl group in poly(p-hydroxystyrene) was replaced by ethoxyethyl groups, and 10 mol% by t-butyl groups. This resin is indicated as a
“resin (B-7)”.

20 [0177]

Synthesis Example 11

A monomer solution was prepared by dissolving 53.69 g of 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate and 46.31 g of 2-methyladamantan-2-yl methacrylate in 200 g of 2-butanone, and further adding 4.04 g
25 of dimethyl azobisisobutyrate.

A 1,000 ml three-necked flask containing 100 g of 2-butanone was purged with nitrogen for 30 minutes and heated to 80°C while stirring. Then, the above monomer

solution was added dropwise to the flask using a dripping funnel over four hours. The polymerization reaction was carried out for six hours after initiation of dripping. After the polymerization, the polymer solution was cooled with water to 30°C or less and poured into 2,000 g of methanol. White precipitate produced was collected by
 5 filtration. The resulting white powder was washed twice with 400 g of methanol in the form of a slurry, filtered, and dried for 17 hours at 50°C to obtain a white resin powder.

The resin was found to have a Mw of 9,700 and confirmed to be a copolymer of 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate and 2-methyladamantan-2-yl methacrylate at a copolymerization molar ratio of 59.6:40.4. This resin is indicated as
 10 a "resin (B-8)".

[0178]

Synthesis Example 12

A monomer solution was prepared by dissolving 40.90 g of 2-methyladamantan-2-yl methacrylate, 15.47 g of 3-hydroxyadamantan-1-yl
 15 methacrylate, and 43.64 g of 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate in 200 g of 2-butanone, and further adding 4.02 g of dimethyl azobisbutyrate.

A 1,000 ml three-necked flask containing 100 g of 2-butanone was purged with nitrogen for 30 minutes and heated to 80°C while stirring. Then, the above monomer solution was added dropwise to the flask using a dripping funnel over four hours. The
 20 polymerization reaction was carried out for six hours after initiation of dripping. After the polymerization, the polymer solution was cooled with water to 30°C or less and poured into 2,000 g of methanol. White precipitate produced was collected by filtration. The resulting white powder was washed twice with 400 g of methanol in the form of a slurry, filtered, and dried for 17 hours at 50°C to obtain a white resin powder.

The resin was found to have a Mw of 9,200 and confirmed to be a copolymer of 2-methyladamantan-2-yl methacrylate, 3-hydroxyadamantan-1-yl methacrylate, and
 25 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate at a copolymerization molar

ratio of 36.2:15.2:48.6. This resin is indicated as a "resin (B-9)".

[0179]

Synthesis Example 13

A monomer solution was prepared by dissolving 43.66 g of
 5 1-(adamantan-1-yl)-1-methylethyl methacrylate, 14.74 g of 3-hydroxyadamantan-1-yl
 methacrylate, and 43.66 g of 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate in
 200 g of 2-butanone, and further adding 3.83 g of dimethyl azobisisobutyrate.

A 1,000 ml three-necked flask containing 100 g of 2-butanone was purged with
 nitrogen for 30 minutes and heated to 80°C while stirring. Then, the above monomer
 10 solution was added dropwise to the flask using a dripping funnel over four hours. The
 polymerization reaction was carried out for six hours after initiation of dripping. After
 the polymerization, the polymer solution was cooled with water to 30°C or less and
 poured into 2,000 g of methanol. White precipitate produced was collected by
 filtration. The resulting white powder was washed twice with 400 g of methanol in the
 15 form of a slurry, filtered, and dried for 17 hours at 50°C to obtain a white resin powder.

The resin was found to have a Mw of 9,600 and confirmed to be a copolymer of
 1-(adamantan-1-yl)-1-methylethyl, 3-hydroxyadamantan-1-yl ester methacrylate, and
 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate at a copolymerization molar
 ratio of 35.6:15.1:49.3. This resin is indicated as a "resin (B-10)".

20 [0180]

Synthesis Example 14

A monomer solution was prepared by dissolving 16.13 g of
 2-ethyladamantan-2-yl methacrylate, 40.58 g of 2-methyladamantan-2-yl methacrylate,
 and 43.29 g of 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate in 200 g of
 25 2-butanone, and further adding 3.99 g of dimethyl azobisisobutyrate.

A 1,000 ml three-necked flask containing 100 g of 2-butanone was purged with
 nitrogen for 30 minutes and heated to 80°C while stirring. Then, the above monomer

solution was added dropwise to the flask using a dripping funnel over four hours. The polymerization reaction was carried out for six hours after initiation of dripping. After the polymerization, the polymer solution was cooled with water to 30°C or less and poured into 2,000 g of methanol. White precipitate produced was collected by

5 filtration. The resulting white powder was washed twice with 400 g of methanol in the form of a slurry, filtered, and dried for 17 hours at 50°C to obtain a white resin powder.

The resin was found to have a Mw of 8,900 and confirmed to be a copolymer of 2-ethyladamantan-2-yl methacrylate, 2-methyladamantan-2-yl methacrylate, and 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate at a copolymerization molar
10 ratio of 13.7:38.2:48.1. This resin is indicated as a "resin (B-11)".

[0181]

Synthesis Example 15

A monomer solution was prepared by dissolving 42.44 g of 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl acrylate, 15.10 g of 3-hydroxyadamantan-1-yl
15 acrylate, and 42.46 g of 2-ethyladamantan-2-yl acrylate in 200 g of 2-butanone, and further adding 4.17 g of dimethyl azobisbutyrate.

A 1,000 ml three-necked flask containing 100 g of 2-butanone was purged with nitrogen for 30 minutes and heated to 80°C while stirring. Then, the above monomer solution was added dropwise to the flask using a dripping funnel over four hours. The
20 polymerization reaction was carried out for six hours after initiation of dripping. After the polymerization, the polymer solution was cooled with water to 30°C or less and poured into 2,000 g of methanol. White precipitate produced was collected by filtration. The resulting white powder was washed twice with 400 g of methanol in the form of a slurry, filtered, and dried for 17 hours at 50°C to obtain a white resin powder.

25 The resin was found to have a Mw of 10,200 and confirmed to be a copolymer of 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl acrylate, 3-hydroxyadamantan-1-yl acrylate, and 2-ethyladamantan-2-yl acrylate at a copolymerization molar ratio of 49.2:15.3:35.5.

This resin is indicated as a "resin (B-12)".

[0182]

Synthesis Example 16

A monomer solution was prepared by dissolving 55.00 g of
 5 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate, 11.70 g of
 3-hydroxyadamantan-1-yl methacrylate, and 33.31 g of 1-ethylcyclopentyl acrylate in
 200 g of 2-butanone, and further adding 4.56 g of dimethyl azobisbutyrate.

A 1,000 ml three-necked flask containing 100 g of 2-butanone was purged with
 nitrogen for 30 minutes and heated to 80°C while stirring. Then, the above monomer
 10 solution was added dropwise to the flask using a dripping funnel over four hours. The
 polymerization reaction was carried out for six hours after initiation of dripping. After
 the polymerization, the polymer solution was cooled with water to 30°C or lower and
 poured into 2,000 g of a mixed solvent of 2-propanol/n-heptane at a weight ratio of 1/2
 in the form of a slurry. White precipitate produced was collected by filtration. The
 15 resulting white powder was washed twice with 400 g of a mixed solvent of
 2-propanol/n-heptane at a weight ratio of 1/2, filtered, and dried for 17 hours at 50°C to
 obtain a white resin powder.

The resin was found to have a Mw of 8,500 and confirmed to be a copolymer of
 5-oxo-4-oxatricyclo[4.2.1.0^{3,7}]nonan-2-yl methacrylate, 3-hydroxyadamantan-1-yl
 20 methacrylate, and 1-ethylcyclopentyl acrylate at a copolymerization molar ratio of
 53.7:11.1:35.2. This resin is indicated as a "resin (B-13)".

[0183]

Synthesis Example 17

A three-necked flask equipped with a stirrer, a cold current condenser, and a
 25 thermometer was charged with 1.52 g of a triethoxysilane compound providing the
 recurring unit of the above formula (12-2), 1.57 g of a triethoxysilane compound
 providing the recurring unit of the above formula (14-1), 1.91 g of

methyltriethoxysilane, 15 g of 4-methyl-2-pentanone, and 1.31 g of a 1.75 wt% aqueous solution of oxalic acid. The mixture was reacted for six hours at 80°C while stirring, followed by cooling to terminate the reaction. The reaction mixture was poured into a separating funnel to remove the water layer. The organic layer was repeatedly washed
5 with ion-exchanged water until the reaction solution becomes neutral. The organic layer was evaporated under reduced pressure.

The resin obtained was found to have a Mw of 2,000 and confirmed to possess the recurring unit of the formula (12-2) and the recurring unit of the formula (14-1) at a molar ratio of 60:40. This resin is indicated as a “resin (B-14)”.

10 [0184]

<Preparation of composition solution>

Components shown in Table 1-1 and Table 1-2 (wherein part(s) indicates part(s) by weight) were mixed to prepare homogeneous solutions. The solutions were filtered through a membrane filter with a pore diameter of 0.2 μm to prepare the solution
15 compositions of Examples 1-16 and Comparative Examples 1-2.

In addition to the acid generators (A-1)-(A-3) and resins (B-1)-(B-14) of the Examples and Comparative Examples, the following components were used.

Other acid generators

a-1: Triphenylsulfonium trifluoromethanesulfonate

20 a-2: Bis(4-t-butylphenyl)iodonium trifluoromethanesulfonate

a-3:

N-(trifluoromethanesulfonyloxy)bicyclo[2.2.1]hept-5-ene-2,3-dicarboxyimide

[0185]

Acid diffusion controller

25 C-1: Tri-n-hexylamine

C-2: Triethanolamine

C-3: 2-Phenylbenzimidazole

C-4: 1,2-dimethylimidazole

C-5: 3-Pyridino-1,2-propanediol

Dissolution controller

D-1: t-Butoxycarbonylmethyl deoxycholate

5 Solvent

S-1: Ethyl lactate

S-2: Propylene glycol monomethyl ether acetate

S-3: 2-heptanone

S-4: γ -Butyrolacton

10 [0186]

<Evaluation>

The solution compositions of Examples 1-16 and Comparative Examples 1-2 were spin-coated on a silicon wafer. PB was then performed under the conditions shown in Table 2 to form resist coatings with a thickness indicated in Table 2. In accordance with the conditions shown in Table 2, exposure and PEB were conducted. The resist patterns were obtained by developing the resist at 23°C for one minute by a paddle method using a 2.38 wt% tetramethylammonium hydroxide aqueous solution, followed by washing with purified water and drying.

[0187]

20 The following were used as the exposure equipment: Stepper NSR2205 EX12B (numerical aperture: 0.55, manufactured by Nikon Corp.) as the KrF excimer laser (indicated by "KrF" in Table 2); ArF excimer laser photolithography machine (numerical aperture: 0.55, manufactured by Nikon Corp.) as the ArF excimer laser (indicated by "ArF" in Table 2); F₂ excimer laser photolithography machine XLS 25 (numerical aperture: 0.60, manufactured by Ultratech) as the F₂ excimer laser; Electron Beam Direct Writing System HL700 (acceleration voltage: 30 KeV, manufactured by Hitachi, Ltd.) improved in acceleration voltage to 50 KeV as the electron beam.

[0188]

Each of the resist patterns obtained was evaluated according to the following method. The evaluation results are shown in Table 3.

Sensitivity

- 5 Sensitivity was evaluated based on an optimum exposure dose which is a dose capable of forming a 1:1 line and space pattern (1L1S) with a line width of 0.25 μm , when a resist film formed on a silicon wafer is exposed to light at a varied dose, immediately followed by PEB, development, washing with water, and drying.

Resolution

- 10 The minimum dimension of a resist pattern resolved at the optimum dose was taken as the resolution of the resist film.

[0189]

Pattern shape

- 15 The dimensions of the lower side L_a and the upper side L_b of the rectangular cross-section of a line and space pattern (1L1S) with a line width of 0.25 μm were measured using a scanning electron microscope. A pattern profile which satisfied a formula " $0.85 \leq L_b/L_a \leq 1$ " was evaluated as "Good", and otherwise evaluated as "Bad".

Storage stability

- 20 The sensitivity of each liquid composition was evaluated immediately after preparation and after standing at room temperature for one month. After standing for one month, the compositions exhibiting a change in sensitivity of less than 10% as compared with immediately after preparation were evaluated as "Good" and those exhibiting a change of 10% or more were evaluated as "Bad".

25

[0190]

[Table 1-1]

	Photoacid generator (Parts by weight)	Acid-dissociable group-containing resin (B) (Parts by weight)	Acid diffusion controller (Parts by weight)	Dissolution controller (Parts by weight)	Solvent (Parts by weight)
Example 1	A-1 (3)	B-5 (65) B-4 (35)	C-2 (0.3)	-	S-1 (250) S-2 (550)
Example 2	A-2 (2) a-3 (2)	B-5 (70) B-3 (30)	C-1 (0.25)	-	S-2 (800)
Example 3	A-1 (2) a-3 (2)	B-7 (100)	C-2 (0.3)	-	S-1 (250) S-2 (550)
Example 4	A-3 (2) a-3 (2)	B-1 (100)	C-4 (0.4)	-	S-1 (800)
Example 5	A-3 (2) a-2(1)	B-2 (100)	C-2 (0.3)	-	S-1 (400) S-2 (400)
Example 6	A-2 (2) a-3 (1)	B-6 (100)	C-2 (0.1) C-4 (0.2)	-	S-1 (250) S-2 (550)
Example 7	A-1 (3)	B-8 (100)	C-5 (0.25)	D-1 (8)	S-2 (550) S-4 (50)
Example 8	A-2 (3)	B-8 (100)	C-2 (0.25)	-	S-2 (550) S-4 (50)
Example 9	A-3 (5)	B-8 (100)	C-2 (0.25)	-	S-2 (550) S-4 (50)
Example 10	A-2 (4)	B-9 (100)	C-5 (0.25)	-	S-2 (550) S-4 (50)

[0191]

5 [Table 1-2]

	Photoacid generator (Parts by weight)	Acid-dissociable group-containing resin (B) (Parts by weight)	Acid diffusion controller (Parts by weight)	Dissolution controller (Parts by weight)	Solvent (Parts by weight)
Example 11	A-2 (4)	B-10 (100)	C-5 (0.25)	-	S-2 (550) S-4 (50)
Example 12	A-2 (4)	B-11 (100)	C-5 (0.25)	D-1 (8)	S-2 (550) S-4 (50)
Example 13	A-2 (4)	B-12 (100)	C-2 (0.25)	-	S-2 (600)
Example 14	A-2 (4)	B-13 (100)	C-2 (0.25)	-	S-2 (550)
Example 15	A-2 (3)	B-1 (100)	C-4 (0.3)	-	S-1 (400) S-2 (400)
Example 16	A-3 (1.5)	B-14 (100)	C-3 (0.1)	-	S-3 (900)
Comparative Example 1	a-1 (5)	B-5 (65) B-4 (35)	C-1 (0.2)	-	S-1 (250) S-2 (550)
Comparative Example 2	a-1 (5)	B-9 (100)	C-1 (0.3)	-	S-1 (400)

[0192]

[Table 2]

	Thickness (Å)	PB		Exposure source	PEB	
		Temperature (°C)	Time (seconds)		Temperature (°C)	Time (seconds)
Example 1	5000	100	90	KrF	110	90
Example 2	5000	100	90	KrF	110	90
Example 3	5000	100	90	KrF	100	90
Example 4	5000	120	90	KrF	130	90
Example 5	5000	120	90	KrF	140	90
Example 6	5000	100	90	KrF	110	90
Example 7	3300	130	90	ArF	130	90
Example 8	3300	130	90	ArF	130	90
Example 9	3300	130	90	ArF	130	90
Example 10	3300	130	90	ArF	130	90
Example 11	3300	130	90	ArF	120	90
Example 12	3300	120	90	ArF	100	90
Example 13	3300	130	90	ArF	100	90
Example 14	3300	130	90	ArF	110	90
Example 15	3000	120	90	Electron beam	130	90
Example 16	1000	130	90	F2	110	90
Comparative Example 1	5000	100	90	KrF	110	90
Comparative Example 2	5000	130	90	ArF	140	90

5

[0193]

[Table 3]

	Sensitivity	Resolution (μm)	Pattern shape	Storage stability
Example 1	320 J/m ²	0.15	Good	Good
Example 2	330 J/m ²	0.15	Good	Good
Example 3	350 J/m ²	0.15	Good	Good
Example 4	350 J/m ²	0.14	Good	Good
Example 5	370 J/m ²	0.16	Good	Good
Example 6	320 J/m ²	0.15	Good	Good
Example 7	382 J/m ²	0.14	Good	Good
Example 8	403 J/m ²	0.14	Good	Good
Example 9	582 J/m ²	0.13	Good	Good
Example 10	443 J/m ²	0.13	Good	Good
Example 11	452 J/m ²	0.13	Good	Good
Example 12	352 J/m ²	0.14	Good	Good
Example 13	337 J/m ²	0.13	Good	Good
Example 14	368 J/m ²	0.13	Good	Good
Example 15	0.03 C/m ²	0.14	Good	Good
Example 16	190 J/m ²	0.13	Good	Good
Comparative Example 1	360 J/m ²	0.16	Bad	Bad
Comparative Example 2	700 J/m ²	0.16	Bad	Bad

[0194]

Table 3 clearly shows that the positive-tone radiation-sensitive resin composition using the acid generator (A) of the present invention excels in storage stability and base resistance, is highly sensitive, and has a high resolution as compared with the positive-tone radiation-sensitive resin compositions of the Comparative Examples.

[0195]

[Effect of the Invention]

The acid generator (A1), acid generator (A2), acid generator (A3), and acid generator (A4) of the present invention excel in storage stability and are highly sensitive to active rays, particularly deep ultraviolet rays represented by a KrF excimer laser, ArF excimer laser, and F₂ excimer laser and electron beams such as EUV. Furthermore, the positive-tone radiation-sensitive resin composition of the present invention using these acid generators excels in storage stability, is highly sensitive, has a high resolution, and is very useful as a chemically amplified resist in the manufacture of integrated circuit

elements of which downsizing is anticipated to proceed in the future. The onium salt compound (2), onium salt compound (3), and onium salt compound (4) of the present invention are highly sensitive to active rays, for example, deep ultraviolet rays such as a KrF excimer laser, ArF excimer laser, and F₂ excimer laser and electron beams such as EUV, and is very useful as a photoacid generator.

[Brief Description of the Drawings]

[Figure 1]

Figure 1 shows a mass spectrometry spectrum in the cation moiety of the acid generator (A-1).

10 [Figure 2]

Figure 2 shows a mass spectrometry spectrum in the cation moiety of the acid generator (A-2).

[Figure 3]

Figure 3 shows a mass spectrometry spectrum in the cation moiety of the acid generator (A-3).

15 [Figure 4]

Figure 4 shows a ¹H-NMR spectrometry spectrum of the acid generator (A-1).

[Figure 5]

Figure 5 shows a ¹H-NMR spectrometry spectrum of the acid generator (A-2).

20 [Figure 6]

Figure 6 shows a ¹H-NMR spectrometry spectrum of the acid generator (A-3).

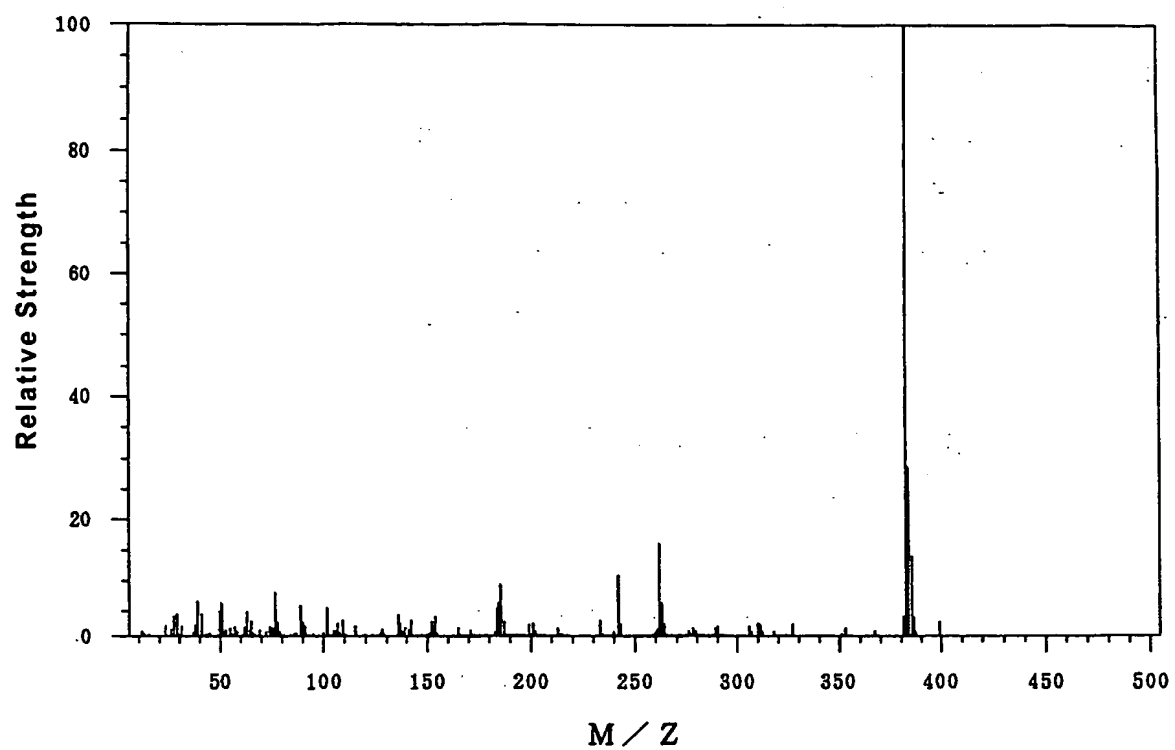


FIG 1

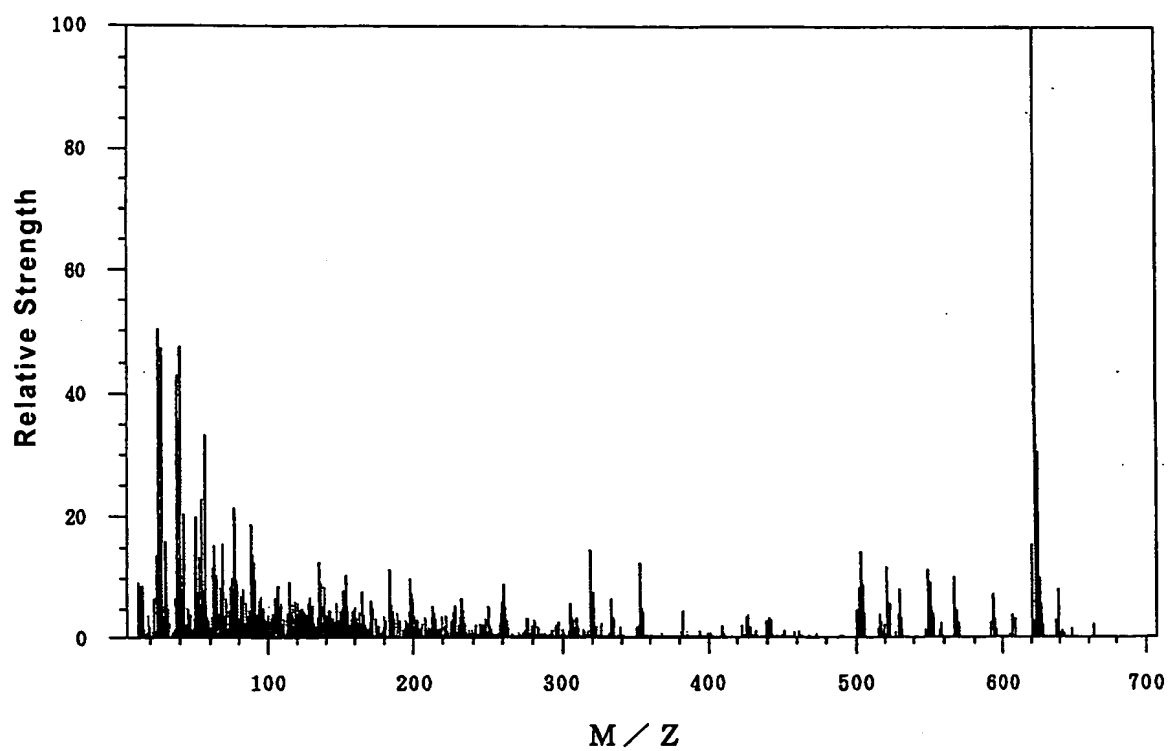


FIG 2

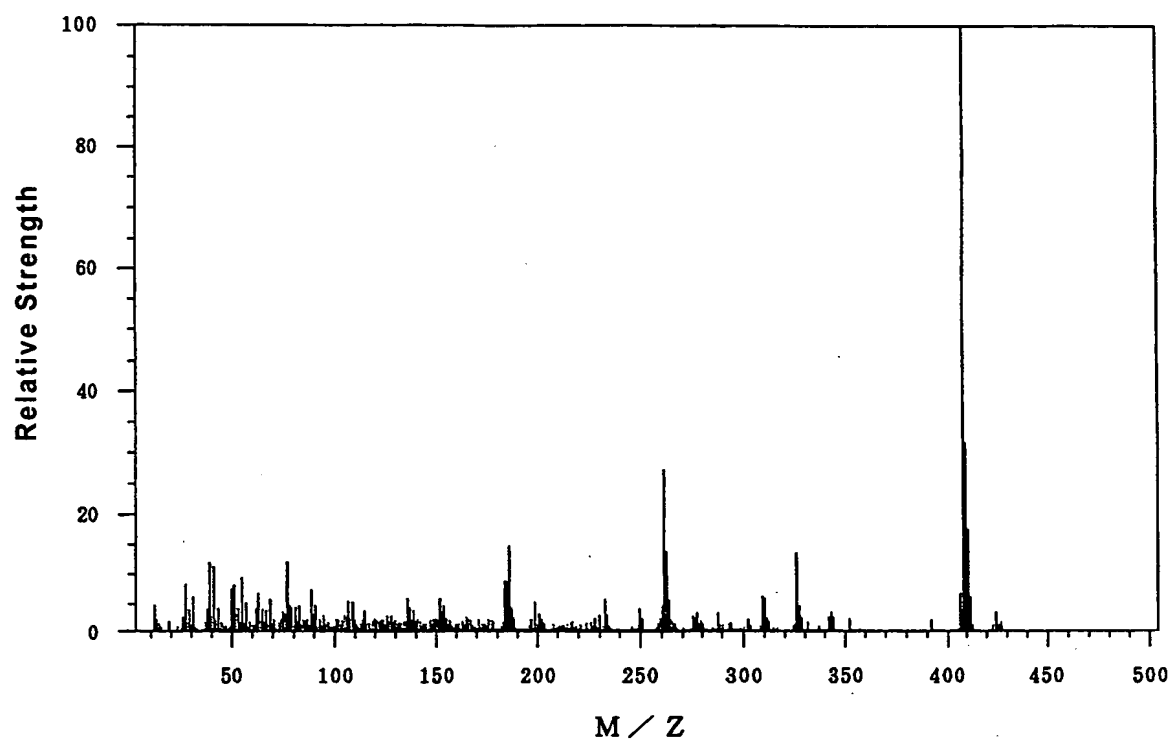


FIG 3

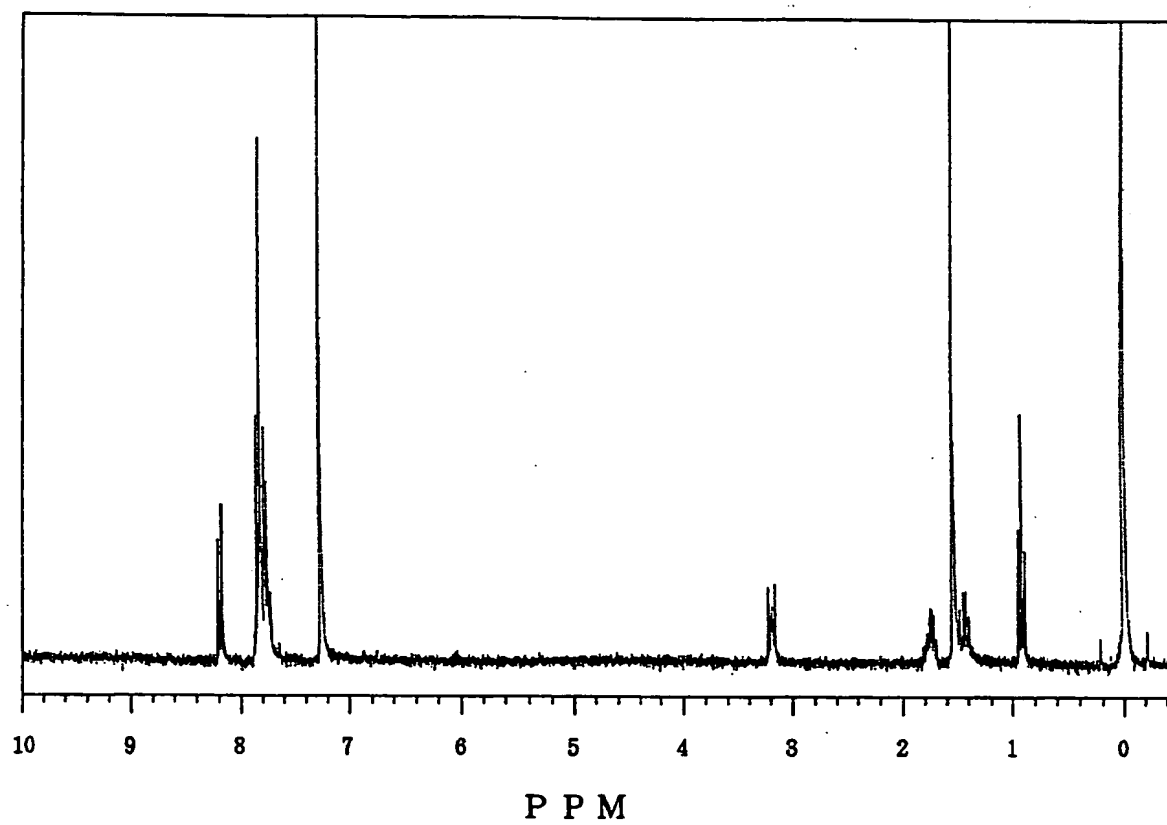


FIG 4

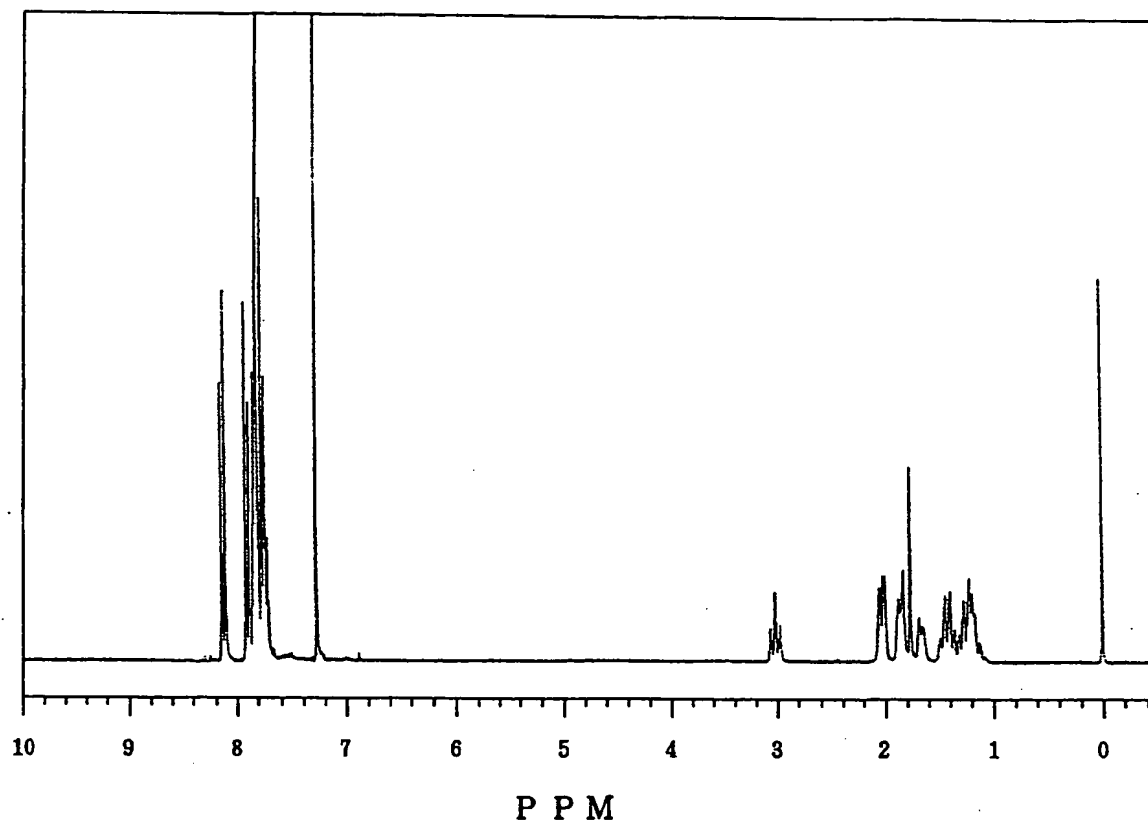


FIG 5

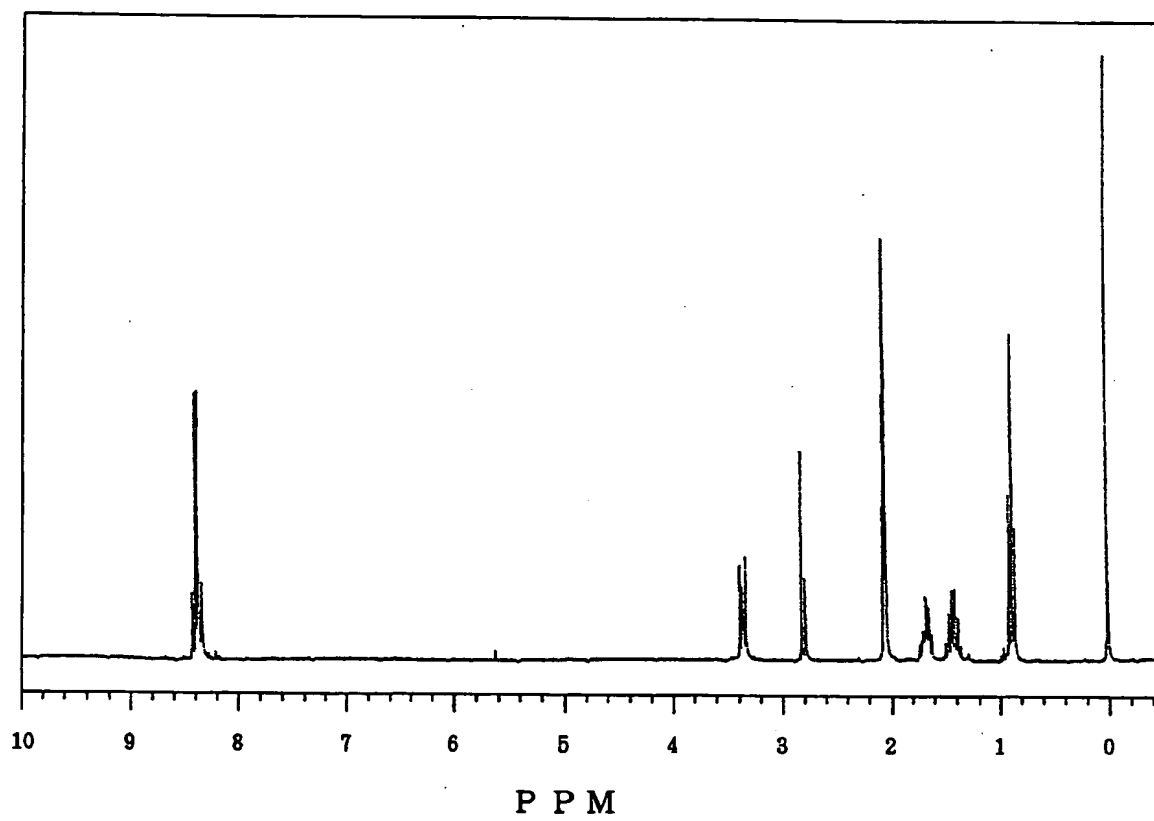


FIG 6

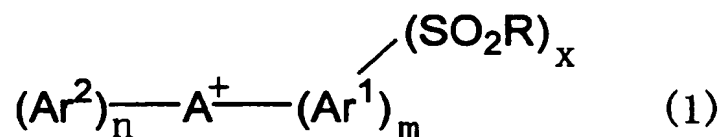
[Document Name] Abstract

[Abstract]

[Object] A photoacid generator (for photoresist) having superior storage stability, used for a radiation-sensitive resin composition with a high sensitivity and high resolution, an onium salt compound suitable for the photoacid generator, and a positive-tone radiation-sensitive resin composition comprising the photoacid generator are provided.

[Means for the Solution] A photoacid generator comprising an onium salt compound having a cation moiety represented by the formula (1),

[Chemical Formula 1]



wherein A represents I or S, m is an integer more than 1, n is an integer more than 0, x is an integer of 1-15, Ar¹ and Ar² represent a (substituted) monovalent aromatic hydrocarbon group and the like, and R represents a (substituted) alkyl group, a (substituted) monovalent alicyclic hydrocarbon group, and the like.

An onium salt compound comprising a compound having a cation moiety represented by the formula (1) in which R is a (substituted) alicyclic hydrocarbon group.

A positive-tone radiation-sensitive resin composition comprising (A) the photoacid generator and (B) an acid-dissociable group-containing resin.

[Selective Drawing] None

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This is to certify that the annexed is a true copy of the following application as filed with this Office.

Date of Application:	December 25, 2002
Application Number:	2002-373625
[ST.10/C]:	JP2002-373625
Applicant(s):	JSR Corporation

December 11, 2003

Commissioner, Patent Office
Yasuo IMAI

Certificate No. 2003-3102674

[Document Name]

APPLICATION FOR PATENT

[Preference Number]

SRR10184

[Filing Date]

December 25, 2002

[Submitted to]

Commissioner, Patent Office

[International Patent Classification]

C07C309/25

C07C309/28

C07C309/71

C07C309/72

G03F 7/004

H01L 21/027

H01L 21/30

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[Designation of Fees]

[Advanced Payment Registration Number]	044428
[Amount Paid]	21,000 Yen

[List of Appended Documents]

[Document Name]	Specification 1
[Document Name]	Drawings 1
[Document Name]	Abstract 1
[General Power of Attorney No.]	9116687

[Proof Requirement]	Requested
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